

PATENT COOPERATION TREATY

DOCKETED

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

MAR 19 2001

To:

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PCT

DUE DATE:

BY:

NOTIFICATION OF TRANSMITTAL OF
THE INTERNATIONAL PRELIMINARY
EXAMINATION REPORT

(PCT Rule 71.1)

Date of mailing
(day/month/year)

08.03.2001

Applicant's or agent's file reference
98,664-A

IMPORTANT NOTIFICATION

International application No.
PCT/US99/26788

International filing date (day/month/year)
10/11/1999

Priority date (day/month/year)
10/11/1998

Applicant

DIGITAL GENE TECHNOLOGIES, INC ET AL

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/

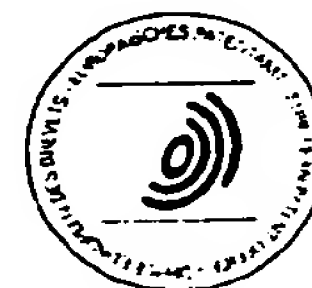


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PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 98,664-A	FOR FURTHER ACTION		See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/US99/26788	International filing date (<i>day/month/year</i>) 10/11/1999	Priority date (<i>day/month/year</i>) 10/11/1998	
International Patent Classification (IPC) or national classification and IPC C12N15/12			
Applicant DIGITAL GENE TECHNOLOGIES, INC ET AL			

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.


2. This REPORT consists of a total of 8 sheets, including this cover sheet.

- ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

- I ☒ Basis of the report
- II ☐ Priority
- III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain documents cited
- VII ☒ Certain defects in the international application
- VIII ☒ Certain observations on the international application

Date of submission of the demand 06/06/2000	Date of completion of this report 08.03.2001
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Chavanne, F Telephone No. +49 89 2399 8399



**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US99/26788

I. Basis of the report

1. This report has been drawn on the basis of *(substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments (Rules 70.16 and 70.17).):*

Description, pages:

1-64 as originally filed

Claims, No.:

1-32 as originally filed

Drawings, sheets:

1/23-23/23 as originally filed

Sequence listing part of the description, pages:

1-19, as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☒ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US99/26788

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

- ☐ the entire international application.
- ☒ claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 4, 6-12, 15-17, 29 (partially).

because:

- ☐ the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (*specify*):
 - ☐ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):
 - ☒ the claims, or said claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 9-12, 15-17 (partially) are so inadequately supported by the description that no meaningful opinion could be formed.
 - ☒ no international search report has been established for the said claims Nos. 1-32 (partially).
2. A meaningful international preliminary examination report cannot be carried out due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:
- ☐ the written form has not been furnished or does not comply with the standard.
 - ☐ the computer readable form has not been furnished or does not comply with the standard.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US99/26788

1. Statement

Novelty (N)	Yes:	Claims	4, 6-8, 29
	No:	Claims	9-12, 15-17
Inventive step (IS)	Yes:	Claims	
	No:	Claims	4, 6-12, 15-17, 29
Industrial applicability (IA)	Yes:	Claims	4, 6-12, 15-17, 29
	No:	Claims	

2. Citations and explanations **see separate sheet**

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. As a result of an objection of lack of unity raised by the International Search Authority, the international search report has been restricted to the invention referred to as invention 1 (claims 1-32, partially), namely:
a polypeptide IMX4 represented by amino acid sequence ID No. 27 and DNA molecules represented by nucleic acid sequences ID No. 1 and ID No. 11, 12 and 13, antibodies binding to said polypeptide, host cells expressing said polypeptide, variants thereof, and therapeutic uses thereof.
Consequently, an opinion with regard to novelty, inventive step and industrial applicability can only be partially formulated, and limited to the above mentioned invention 1.
2. An opinion with regard to novelty, inventive step and industrial applicability cannot be formulated for the claims relating to a polypeptide comprising the amino acid sequence of SEQ ID No. 27, or a polypeptide encoded by a polynucleotide of SEQ ID No. 1, for the following reasons:

According to the present application, the polypeptide having the amino acid sequence of SEQ ID No. 27 is the translational product of the nucleotide sequence of isolated DNA molecule IMX4 of SEQ ID No. 1 (see e.g. page 59, example 2). However, in view of the nucleotide sequence of SEQ ID No. 1, it appears that said DNA molecule cannot encode said polypeptide:

A. The nucleotide sequence of SEQ ID No. 1 contains only one putative methionine initiation codon (ATG) at position 29. The corresponding putative open reading frame (ORF) stops after three amino acids (KAV) with a stop codon (TAG) at position 41. This is too short to corresponds to an ORF leading to a translated product. Thus, there are at the nucleotide level no indications that the nucleotide sequence of SEQ ID No. 1 encodes for a polypeptide.

B. The comparison of the amino acid sequence of SEQ ID No. 27 with the nucleotide sequence of SEQ ID No. 1 shows that the codon supposed to correspond to the methionine of SEQ ID No. 27 actually corresponds to a stop

codon (TAG). Thus, the polypeptide of SEQ ID No. 27 cannot be translated from IMX4 as asserted in the present application (see e.g. page 59, line 15).

C. The alignment of the nucleotide sequence of SEQ ID No. 1 with the nucleotide sequence of the soares ovary tumour HbHOT Homo sapiens cDNA clone (D1) shows that the nucleotide sequence of SEQ ID No. 1 corresponds to a 3' untranslated region. In fact, the nucleotide sequence of SEQ ID No. 1 shows over 96% identity with the 3' untranslated region of said human cDNA clone. This alignment shows that the last five A nucleotides of SEQ ID No. 1 are part of the polyA tail. A polyadenylation signal motif can also be found at position 114. Moreover, because of the high mutation rate of non-coding sequences, 3' untranslated regions are gene specific. Thus, it can be totally excluded that a 3' untranslated region would show such a high identity with a coding region.

Therefore, the nucleotide sequence of SEQ ID No. 1, which is supposed to encode the polypeptide having the amino acid sequence of SEQ ID No. 27, actually corresponds to a 3' untranslated region.

Since the polypeptide of SEQ ID No. 27 does not exist, and since the nucleotide sequence of SEQ ID No. 1 does not encode any polypeptide, an opinion with regard to novelty, inventive step and industrial applicability on the claims referring to said polypeptides, and related claims cannot be formulated (claims 1-3, 5, 13, 14, 18-28 and 30-32, completely; 9-12 and 15-17, partially).

V. Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

D1: EMBL Database entry HS1272990
Accession Number AA480830
D2: EMBL Database entry HS6802
Accession Number Z82215.1

2. D1 describes the nucleotide sequence of a cDNA clone comprising a sequence showing over 96% identity with SEQ ID No. 1.
D2 describes a nucleotide sequence showing over 97% identity with the nucleotide sequence of SEQ ID No. 1.
Thus, in view of D1 or D2, the subject-matter of claims 9-12 and 15-17 is not novel. Therefore, claims 9-12 and 15-17 do not meet the requirements of Article 33(2) PCT.
3. The nucleotide sequence of SEQ ID No. 1 described in the present application corresponds to the 3' untranslated region of an unknown gene. The present application shows that the transcription of said gene is increased when cells are treated with interferon- γ (example 1). However, this does not give any evidence on the specific function of the product of said gene. The function of the product of the gene corresponding to the 3' untranslated region of nucleotide sequence of SEQ ID No. 1 being unknown, the isolation of said nucleotide does not solve any problem. Since there is no problem to be solved, there cannot be any inventive step involved. Thus, the subject-matter of claims 4, 6-12, 15-17 and 29 is not inventive. Therefore, claims 4, 6-12, 15-17 and 29 do not meet the requirements of Article 33(3) PCT.

VII. Certain defects in the international application

1. Point (d) of claim 9 is lacking.
2. The specification mentions that the sequences of SEQ ID No. 11 and 12 include the sequence of SEQ ID No. 1 (page 59, lines 7-10). It appears that this is not the case.
3. The specification does not define the SEQ ID No. 18, 19 and 21-26 although they are mentioned in the claims.
4. Claim 16 refers to a method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 9. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a

subject. Both claims do not mention how said methods are to be performed.

VIII. Certain observations on the international application

1. The formulation "...comprises..." in claims 4 and 8-12 does not clearly define the scope of the claim. Thus, this expression should be replaced with "consists of" (Article 6 PCT).
2. Claims 9 and 12 lack clarity due to the formulation "hybridisable to". This formulation is vague and imprecise, it does not provide any technical information about the hybridisation conditions, and as a consequence is not adapted to clearly define the scope of the claim. In fact, any nucleic acid molecule may hybridise to the nucleic acid molecule of SEQ ID No. 1, when the hybridisation conditions are adapted to their degree of identity (Article 6 PCT).
3. Claim 1 mentions that the DNA molecule of SEQ ID No. 1 has altered expression in a T84 model of gut barrier function. However, the specification does not provide any evidence for said altered expression. The present application only shows that treatment of the T84 cells with interferon- γ induces expression of the DNA molecule of SEQ ID No. 1. Thus, claim 1 is not based on the description (Article 6-support PCT).
4. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject by determining the presence or absence of a polynucleotide. Apart from vague declarations, the present application does not provide any evidence showing that the irritable bowel disease may be evidenced by determining the presence or absence of the polynucleotide of SEQ ID No. 1. Thus, claim 29 is not based on the description (Article 6-support PCT). Moreover, since a polynucleotide is always either present or absent, the man skilled in the art, faced with the problem of establishing such a diagnosis would not be able to perform it with said method (Article 5 PCT).

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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 98,664-A ,	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/US99/26788	International filing date (day/month/year) 10/11/1999	Priority date (day/month/year) 10/11/1998
International Patent Classification (IPC) or national classification and IPC C12N15/12		
Applicant DIGITAL GENE TECHNOLOGIES, INC ET AL		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.



2. This REPORT consists of a total of 8 sheets, including this cover sheet.

- ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

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3. This report contains indications relating to the following items:

- I ☒ Basis of the report
- II ☐ Priority
- III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain documents cited
- VII ☒ Certain defects in the international application
- VIII ☒ Certain observations on the international application

Date of submission of the demand 06/06/2000	Date of completion of this report 08.03.2001
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Chavanne, F Telephone No. +49 89 2399 8399 

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US99/26788

I. Basis of the report

1. This report has been drawn on the basis of *(substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments (Rules 70.16 and 70.17).):*

Description, pages:

1-64 as originally filed

Claims, No.:

1-32 as originally filed

Drawings, sheets:

1/23-23/23 as originally filed

Sequence listing part of the description, pages:

1-19, as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☒ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US99/26788

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

- ☐ the entire international application.
- ☒ claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 4, 6-12, 15-17, 29 (partially).

because:

- ☐ the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (*specify*):

- ☐ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):

- ☒ the claims, or said claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 9-12, 15-17 (partially) are so inadequately supported by the description that no meaningful opinion could be formed.

- ☒ no international search report has been established for the said claims Nos. 1-32 (partially).

2. A meaningful international preliminary examination report cannot be carried out due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:

- ☐ the written form has not been furnished or does not comply with the standard.
- ☐ the computer readable form has not been furnished or does not comply with the standard.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US99/26788

1. Statement

Novelty (N)	Yes:	Claims	4, 6-8, 29
	No:	Claims	9-12, 15-17
Inventive step (IS)	Yes:	Claims	
	No:	Claims	4, 6-12, 15-17, 29
Industrial applicability (IA)	Yes:	Claims	4, 6-12, 15-17, 29
	No:	Claims	

2. Citations and explanations see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. As a result of an objection of lack of unity raised by the International Search Authority, the international search report has been restricted to the invention referred to as invention 1 (claims 1-32, partially), namely:
a polypeptide IMX4 represented by amino acid sequence ID No. 27 and DNA molecules represented by nucleic acid sequences ID No. 1 and ID No. 11, 12 and 13, antibodies binding to said polypeptide, host cells expressing said polypeptide, variants thereof, and therapeutic uses thereof.
Consequently, an opinion with regard to novelty, inventive step and industrial applicability can only be partially formulated, and limited to the above mentioned invention 1.
2. An opinion with regard to novelty, inventive step and industrial applicability cannot be formulated for the claims relating to a polypeptide comprising the amino acid sequence of SEQ ID No. 27, or a polypeptide encoded by a polynucleotide of SEQ ID No. 1, for the following reasons:

According to the present application, the polypeptide having the amino acid sequence of SEQ ID No. 27 is the translational product of the nucleotide sequence of isolated DNA molecule IMX4 of SEQ ID No. 1 (see e.g. page 59, example 2). However, in view of the nucleotide sequence of SEQ ID No. 1, it appears that said DNA molecule cannot encode said polypeptide:

- A. The nucleotide sequence of SEQ ID No. 1 contains only one putative methionine initiation codon (ATG) at position 29. The corresponding putative open reading frame (ORF) stops after three amino acids (KAV) with a stop codon (TAG) at position 41. This is too short to corresponds to an ORF leading to a translated product. Thus, there are at the nucleotide level no indications that the nucleotide sequence of SEQ ID No. 1 encodes for a polypeptide.
- B. The comparison of the amino acid sequence of SEQ ID No. 27 with the nucleotide sequence of SEQ ID No. 1 shows that the codon supposed to correspond to the methionine of SEQ ID No. 27 actually corresponds to a stop

codon (TAG). Thus, the polypeptide of SEQ ID No. 27 cannot be translated from IMX4 as asserted in the present application (see e.g. page 59, line 15).

C. The alignment of the nucleotide sequence of SEQ ID No. 1 with the nucleotide sequence of the soares ovary tumour HbHOT Homo sapiens cDNA clone (D1) shows that the nucleotide sequence of SEQ ID No. 1 corresponds to a 3' untranslated region. In fact, the nucleotide sequence of SEQ ID No. 1 shows over 96% identity with the 3' untranslated region of said human cDNA clone. This alignment shows that the last five A nucleotides of SEQ ID No. 1 are part of the polyA tail. A polyadenylation signal motif can also be found at position 114. Moreover, because of the high mutation rate of non-coding sequences, 3' untranslated regions are gene specific. Thus, it can be totally excluded that a 3' untranslated region would show such a high identity with a coding region.

Therefore, the nucleotide sequence of SEQ ID No. 1, which is supposed to encode the polypeptide having the amino acid sequence of SEQ ID No. 27, actually corresponds to a 3' untranslated region.

Since the polypeptide of SEQ ID No. 27 does not exist, and since the nucleotide sequence of SEQ ID No. 1 does not encode any polypeptide, an opinion with regard to novelty, inventive step and industrial applicability on the claims referring to said polypeptides, and related claims cannot be formulated (claims 1-3, 5, 13, 14, 18-28 and 30-32, completely; 9-12 and 15-17, partially).

V. Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

- D1: EMBL Database entry HS1272990
Accession Number AA480830
- D2: EMBL Database entry HS6802
Accession Number Z82215.1

2. D1 describes the nucleotide sequence of a cDNA clone comprising a sequence showing over 96% identity with SEQ ID No. 1.
D2 describes a nucleotide sequence showing over 97% identity with the nucleotide sequence of SEQ ID No. 1.
Thus, in view of D1 or D2, the subject-matter of claims 9-12 and 15-17 is not novel. Therefore, claims 9-12 and 15-17 do not meet the requirements of Article 33(2) PCT.
3. The nucleotide sequence of SEQ ID No. 1 described in the present application corresponds to the 3' untranslated region of an unknown gene. The present application shows that the transcription of said gene is increased when cells are treated with interferon- γ (example 1). However, this does not give any evidence on the specific function of the product of said gene. The function of the product of the gene corresponding to the 3' untranslated region of nucleotide sequence of SEQ ID No.1 being unknown, the isolation of said nucleotide does not solve any problem. Since there is no problem to be solved, there cannot be any inventive step involved. Thus, the subject-matter of claims 4, 6-12, 15-17 and 29 is not inventive. Therefore, claims 4, 6-12, 15-17 and 29 do not meet the requirements of Article 33(3) PCT.

VII. Certain defects in the international application

1. Point (d) of claim 9 is lacking.
2. The specification mentions that the sequences of SEQ ID No. 11 and 12 include the sequence of SEQ ID No. 1 (page 59, lines 7-10). It appears that this is not the case.
3. The specification does not define the SEQ ID No. 18, 19 and 21-26 although they are mentioned in the claims.
4. Claim 16 refers to a method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 9. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a

subject. Both claims do not mention how said methods are to be performed.

VIII. Certain observations on the international application

1. The formulation "...comprises..." in claims 4 and 8-12 does not clearly define the scope of the claim. Thus, this expression should be replaced with "consists of" (Article 6 PCT).
2. Claims 9 and 12 lack clarity due to the formulation "hybridisable to". This formulation is vague and imprecise, it does not provide any technical information about the hybridisation conditions, and as a consequence is not adapted to clearly define the scope of the claim. In fact, any nucleic acid molecule may hybridise to the nucleic acid molecule of SEQ ID No. 1, when the hybridisation conditions are adapted to their degree of identity (Article 6 PCT).
3. Claim 1 mentions that the DNA molecule of SEQ ID No. 1 has altered expression in a T84 model of gut barrier function. However, the specification does not provide any evidence for said altered expression. The present application only shows that treatment of the T84 cells with interferon- γ induces expression of the DNA molecule of SEQ ID No. 1. Thus, claim 1 is not based on the description (Article 6-support PCT).
4. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject by determining the presence or absence of a polynucleotide. Apart from vague declarations, the present application does not provide any evidence showing that the irritable bowel disease may be evidenced by determining the presence or absence of the polynucleotide of SEQ ID No. 1. Thus, claim 29 is not based on the description (Article 6-support PCT). Moreover, since a polynucleotide is always either present or absent, the man skilled in the art, faced with the problem of establishing such a diagnosis would not be able to perform it with said method (Article 5 PCT).

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 98,664-A	FOR FURTHER ACTION		see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
International application No. PCT/US 99/ 26788	International filing date (day/month/year) 10/11/1999	(Earliest) Priority Date (day/month/year) 10/11/1998	
Applicant DIGITAL GENE TECHNOLOGIES, INC ET AL			

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 9 sheets.

☐ It is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☒ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☐ furnished subsequently to this Authority in written form.

☒ furnished subsequently to this Authority in computer readable form.

☒ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☒ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☒ **Certain claims were found unsearchable** (See Box I).

3. ☒ **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,

☐ the text is approved as submitted by the applicant.

☒ the text has been established by this Authority to read as follows:

NUCLEIC ACIDS AND POLYPEPTIDES HAVING ALTERED EXPRESSION IN THE T84 GUT BARRIER MODEL

5. With regard to the **abstract**,

☐ the text is approved as submitted by the applicant.

☒ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. _____

☐ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

☒ None of the figures.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 99/26788

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☒ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-32 all partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 29 as far as it concerns an in vivo method is directed to a diagnostic method practised on the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Although claims 27 and 28 are directed to a method of treatment of the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box I.2

It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-32 all partially

A polypeptide IMX4 represented by amino acid sequence ID no.27 and DNA molecules represented by nucleic acid sequences ID no.1 and ID no.11,12 and 13. Host cell. Antibody. Variants thereof. Therapeutic uses.

2. Claims: 1-32 all partially

A polypeptide IMX10 represented by amino acid sequence ID no.28 and DNA molecules represented by nucleic acid sequences ID no.2 and ID no.14. Host cell. Antibody. Variants thereof. Therapeutic uses.

3. Claims: 1-32 all partially

A polypeptide IMX21 represented by amino acid sequence ID no.29 and DNA molecules represented by nucleic acid sequences ID no.3 and ID no.15. Host cell. Antibody. Variants thereof. Therapeutic uses.

4. Claims: 1-32 all partially

A polypeptide IMX28 represented by amino acid sequence ID no.30 and DNA molecules represented by nucleic acid sequences ID no.4 and no.16. Host cell. Antibody. Variants thereof. Therapeutic uses.

5. Claims: 1-32 all partially

A polypeptide IMX32 represented by amino acid sequence ID no.31 and DNA molecules represented by nucleic acid sequences ID no.5, no.17 and no.18. Host cell. Antibody. Variants thereof. Therapeutic uses.

6. Claims: 1-32 all partially

A polypeptide IMX39 represented by amino acid sequence ID no.33 and DNA molecules represented by nucleic acid sequences ID no.20 and no.6. Host cell. Antibody. Variants thereof. Therapeutic uses.

7. Claims: 1-32 all partially

A polypeptide IMX40 represented by amino acid sequence ID no.34 and DNA molecules represented by nucleic acid sequences ID no.7 and ID no.21. Host cell. Antibody. Variants thereof. Therapeutic uses.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

8. Claims: 1-32 all partially

A polypeptide IMX42 represented by amino acid sequence ID no.35 and DNA molecules represented by nucleic acid sequences ID no.8. Host cell. Antibody. Variants thereof. Therapeutic uses.

9. Claims: 1-32 all partially

A polypeptide IMX44 represented by amino acid sequence ID no.37 and DNA molecules represented by nucleic acid sequences ID no.9, no.23 and no.24. Host cell. Antibody. Variants thereof. Therapeutic uses.

10. Claims: 1-32 all partially

A polypeptide IMX56 represented by amino acid sequence ID no.38 and DNA molecules represented by nucleic acid sequences ID no.10 and no.25. Host cell. Antibody. Variants thereof. Therapeutic uses.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/26788

Box III TEXT OF THE ABSTRACT (Continuation of item 5 of the first sheet)

The invention is directed to purified and isolated polypeptides, the nucleic acids encoding such polypeptides that have altered expression characteristics in the T84 gut barrier model, processes for production of recombinant forms of such polypeptides, antibodies generated against these polypeptides, fragmented peptides derived from these polypeptides and the uses to study inflammatory bowel diseases (IBD).

INTERNATIONAL SEARCH REPORT

International Application No

PC 99/26788

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12N15/12 C12Q1/68 A61K38/17 C07K14/47 G01N33/68
C07K16/18 A61P1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>L. HILLIER ET AL: "washU-Merck EST project 1997. zx87d07.r1 Soares tumor NbHOT Homo sapiens cDNA clone IMAGE:810733 5', mRNA sequence" EMBL DATABASE ENTRY HS1272990, ACCESSION NUMBER AA480830;24-June 1997 (1997-06-24), XP002136972 & UNPUBLISHED, (97% identity with the entire seq ID no.1) --- -/-</p>	4,8,9, 12,15-17

☒ Further documents are listed in the continuation of box C.☐ Patent family members are listed in annex.

° Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

4 May 2000

Date of mailing of the international search report

16. 08. 2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

LE CORNEC N.D.R.

INTERNATIONAL SEARCH REPORT

International Application No

PC 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>P.N. DUCHATEAU ET AL: "Homo sapiens apolipoprotein L mRNA, complete cds" EMBL DATABASE ENTRY AF019225, ACCESSION NUMBER AF019225, 29 September 1997 (1997-09-29), XP002136973 cited in the application -& P.N. DUCHATEAU ET AL: "Apolipoprotein L, a new human high density lipoprotein apolipoprotein expressed by the pancreas. Identification, cloning, characterization, and plasma distribution" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 272, no. 41, 1997, pages 25576-25582, XP002136974 AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD., US ISSN: 0021-9258</p> <p>---</p>	8-17
X	<p>D. LLOYD: "Human DNA sequence from clone RP1-6802 on chromosome 22 contains the 5' end of the APOL2 gene for apolipoprotein L2, the APOL gene for apolipoprotein L, the MYH9 for nonmuscle type myosin heavy chain 9. ESTs, STSs and GSSs" EMBL DATABASE ENTRY HS6802, ACCESSION NUMBER Z82215, 8 November 1996 (1996-11-08), XP002136975 " 99,1% identity with whole sequence ID no.11. Reverse orientation" abstract</p> <p>---</p>	8-17
A	<p>F. BESANÇON ET AL: "Interferon-gamma downregulates CFTR gene expression in epithelial cells" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 267, no. 5, November 1994 (1994-11), pages C1398-C1404, XP000906853 the whole document</p> <p>---</p>	1-32
A	<p>J.L. MADARA ET AL: "Interferon-gamma directly affects barrier function of cultured epithelial monolayers" THE JOURNAL OF CLINICAL INVESTIGATION, vol. 83, no. 2, February 1989 (1989-02), pages 724-727, XP000906849 cited in the application the whole document</p> <p>---</p>	1-32

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INTERNATIONAL SEARCH REPORT

International Application No

PO 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	K.DHARMASATHAPHORN ET AL: "A human colonic tumor cell line that maintains vectorial electrolyte transport" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 9, no. 2, February 1984 (1984-02), pages G204-G208, XP002136976 cited in the application ---	
A	J.L. MADARA ET AL: "Occluding junction structure-function relationships in a cultured epithelial monolayer" THE JOURNAL OF CELL BIOLOGY, vol. 101, December 1985 (1985-12), pages 2124-2133, XP000906833 cited in the application ---	
T	A. YOUAKIM ET AL: "Interferon-gamma decreases barrier function in T84 cells by reducing ZO-1 levels and disrupting apical actin" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 276, no. 5, May 1999 (1999-05), pages G1279-G1288, XP000906855 -----	

INTERNATIONAL SEARCH REPORT

National Application No
PCT/US 99/26788

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12N15/12 C12Q1/68 A61K38/17 C07K14/47 G01N33/68
C07K16/18 A61P1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. HILLIER ET AL: "washU-Merck EST project 1997. zx87d07.r1 Soares tumor NbHOT Homo sapiens cDNA clone IMAGE:810733 5', mRNA sequence" EMBL DATABASE ENTRY HS1272990, ACCESSION NUMBER AA480830, 24 June 1997 (1997-06-24), XP002136972 & UNPUBLISHED, (97% identity with the entire seq ID no.1) --- -/--	4,8,9, 12,15-17

☒ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *Z* document member of the same patent family

Date of the actual completion of the international search

4 May 2000

Date of mailing of the international search report

16 08 2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

LE CORNEC N.D.R.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>P.N. DUCHATEAU ET AL: "Homo sapiens apolipoprotein L mRNA, complete cds" EMBL DATABASE ENTRY AF019225, ACCESSION NUMBER AF019225, 29 September 1997 (1997-09-29), XP002136973 cited in the application -& P.N. DUCHATEAU ET AL: "Apolipoprotein L, a new human high density lipoprotein apolipoprotein expressed by the pancreas. Identification, cloning, characterization, and plasma distribution" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 272, no. 41, 1997, pages 25576-25582, XP002136974 AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD., US ISSN: 0021-9258</p> <p style="text-align: center;">---</p>	8-17
X	<p>D. LLOYD: "Human DNA sequence from clone RP1-6802 on chromosome 22 contains the 5' end of the APOL2 gene for apolipoprotein L2, the APOL gene for apolipoprotein L, the MYH9 for nonmuscle type myosin heavy chain 9. ESTs, STSs and GSSs" EMBL DATABASE ENTRY HS6802, ACCESSION NUMBER Z82215, 8 November 1996 (1996-11-08), XP002136975 " 99,1% identity with whole sequence ID no.11. Reverse orientation" abstract</p> <p style="text-align: center;">---</p>	8-17
A	<p>F. BESANCON ET AL: "Interferon-gamma downregulates CFTR gene expression in epithelial cells" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 267, no. 5, November 1994 (1994-11), pages C1398-C1404, XP000906853 the whole document</p> <p style="text-align: center;">---</p>	1-32
A	<p>J.L. MADARA ET AL: "Interferon-gamma directly affects barrier function of cultured epithelial monolayers" THE JOURNAL OF CLINICAL INVESTIGATION, vol. 83, no. 2, February 1989 (1989-02), pages 724-727, XP000906849 cited in the application the whole document</p> <p style="text-align: center;">---</p>	1-32

-/--

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	K.DHARMASATHAPHORN ET AL: "A human colonic tumor cell line that maintains vectorial electrolyte transport" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 9, no. 2, February 1984 (1984-02), pages G204-G208, XP002136976 cited in the application	
A	--- J.L. MADARA ET AL: "Occluding junction structure-function relationships in a cultured epithelial monolayer" THE JOURNAL OF CELL BIOLOGY, vol. 101, December 1985 (1985-12), pages 2124-2133, XP000906833 cited in the application	
T	--- A. YOUAKIM ET AL: "Interferon-gamma decreases barrier function in T84 cells by reducing ZO-1 levels and disrupting apical actin" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 276, no. 5, May 1999 (1999-05), pages G1279-G1288, XP000906855 -----	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 99/26788

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☒ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-32 all partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 29 as far as it concerns an in vivo method is directed to a diagnostic method practised on the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Although claims 27 and 28 are directed to a method of treatment of the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box I.2

It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-32 all partially

A polypeptide IMX4 represented by amino acid sequence ID no.27 and DNA molecules represented by nucleic acid sequences ID no.1 and ID no.11,12 and 13. Host cell. Antibody. Variants thereof. Therapeutic uses.

2. Claims: 1-32 all partially

A polypeptide IMX10 represented by amino acid sequence ID no.28 and DNA molecules represented by nucleic acid sequences ID no.2 and ID no.14. Host cell. Antibody. Variants thereof. Therapeutic uses.

3. Claims: 1-32 all partially

A polypeptide IMX21 represented by amino acid sequence ID no.29 and DNA molecules represented by nucleic acid sequences ID no.3 and ID no.15. Host cell. Antibody. Variants thereof. Therapeutic uses.

4. Claims: 1-32 all partially

A polypeptide IMX28 represented by amino acid sequence ID no.30 and DNA molecules represented by nucleic acid sequences ID no.4 and no.16. Host cell. Antibody. Variants thereof. Therapeutic uses.

5. Claims: 1-32 all partially

A polypeptide IMX32 represented by amino acid sequence ID no.31 and DNA molecules represented by nucleic acid sequences ID no.5, no.17 and no.18. Host cell. Antibody. Variants thereof. Therapeutic uses.

6. Claims: 1-32 all partially

A polypeptide IMX39 represented by amino acid sequence ID no.33 and DNA molecules represented by nucleic acid sequences ID no.20 and no.6. Host cell. Antibody. Variants thereof. Therapeutic uses.

7. Claims: 1-32 all partially

A polypeptide IMX40 represented by amino acid sequence ID no.34 and DNA molecules represented by nucleic acid sequences ID no.7 and ID no.21. Host cell. Antibody. Variants thereof. Therapeutic uses.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

8. Claims: 1-32 all partially

A polypeptide IMX42 represented by amino acid sequence ID no.35 and DNA molecules represented by nucleic acid sequences ID no.8. Host cell. Antibody. Variants thereof. Therapeutic uses.

9. Claims: 1-32 all partially

A polypeptide IMX44 represented by amino acid sequence ID no.37 and DNA molecules represented by nucleic acid sequences ID no.9, no.23 and no.24. Host cell. Antibody. Variants thereof. Therapeutic uses.

10. Claims: 1-32 all partially

A polypeptide IMX56 represented by amino acid sequence ID no.38 and DNA molecules represented by nucleic acid sequences ID no.10 and no.25. Host cell. Antibody. Variants thereof. Therapeutic uses.

PATENT COOPERATION TREATY

AUG 23 2000

From the INTERNATIONAL SEARCHING AUTHORITY

PCT

NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL SEARCH REPORT OR THE DECLARATION

(PCT Rule 44.1)

To:

**McDONNELL BOEHNEN HULBERT
& BERGHOFF**
Attn. Zimmerman, Roger P.
300 South Wacker Drive, Suite 3200
CHICAGO, IL 60606
UNITED STATES OF AMERICA

Date of mailing
(day/month/year)

16/08/2000

Applicant's or agent's file reference

98,664-A

FOR FURTHER ACTION

See paragraphs 1 and 4 below

International application No.

PCT/US 99/26788

International filing date
(day/month/year)

10/11/1999

Applicant

DIGITAL GENE TECHNOLOGIES, INC ET AL

1. ☒ The applicant is hereby notified that the International Search Report has been established and is transmitted herewith.

Filing of amendments and statement under Article 19:

The applicant is entitled, if he so wishes, to amend the claims of the International Application (see Rule 46):

When? The time limit for filing such amendments is normally 2 months from the date of transmittal of the International Search Report; however, for more details, see the notes on the accompanying sheet.

Where? Directly to the International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland
Facsimile No.: (41-22) 740.14.35

For more detailed instructions, see the notes on the accompanying sheet.

2. ☐ The applicant is hereby notified that no International Search Report will be established and that the declaration under Article 17(2)(a) to that effect is transmitted herewith.

3. ☐ With regard to the protest against payment of (an) additional fee(s) under Rule 40.2, the applicant is notified that:

☐ the protest together with the decision thereon has been transmitted to the International Bureau together with the applicant's request to forward the texts of both the protest and the decision thereon to the designated Offices.

☐ no decision has been made yet on the protest; the applicant will be notified as soon as a decision is made.

4. **Further action(s):** The applicant is reminded of the following:

Shortly after 18 months from the priority date, the international application will be published by the International Bureau. If the applicant wishes to avoid or postpone publication, a notice of withdrawal of the international application, or of the priority claim, must reach the International Bureau as provided in Rules 90bis.1 and 90bis.3, respectively, before the completion of the technical preparations for international publication.

Within 19 months from the priority date, a demand for international preliminary examination must be filed if the applicant wishes to postpone the entry into the national phase until 30 months from the priority date (in some Offices even later).

Within 20 months from the priority date, the applicant must perform the prescribed acts for entry into the national phase before all designated Offices which have not been elected in the demand or in a later election within 19 months from the priority date or could not be elected because they are not bound by Chapter II.

Name and mailing address of the International Searching Authority



European Patent Office, P.B. 5818 Patentlaan 2
NL-2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Catherine Humbert

NOTES TO FORM PCT/ISA/220

These Notes are intended to give the basic instructions concerning the filing of amendments under article 19. The Notes are based on the requirements of the Patent Cooperation Treaty, the Regulations and the Administrative Instructions under that Treaty. In case of discrepancy between these Notes and those requirements, the latter are applicable. For more detailed information, see also the PCT Applicant's Guide, a publication of WIPO.

In these Notes, "Article", "Rule", and "Section" refer to the provisions of the PCT, the PCT Regulations and the PCT Administrative Instructions, respectively.

INSTRUCTIONS CONCERNING AMENDMENTS UNDER ARTICLE 19

The applicant has, after having received the international search report, one opportunity to amend the claims of the international application. It should however be emphasized that, since all parts of the international application (claims, description and drawings) may be amended during the international preliminary examination procedure, there is usually no need to file amendments of the claims under Article 19 except where, e.g. the applicant wants the latter to be published for the purposes of provisional protection or has another reason for amending the claims before international publication. Furthermore, it should be emphasized that provisional protection is available in some States only.

What parts of the international application may be amended?

Under Article 19, only the claims may be amended.

During the international phase, the claims may also be amended (or further amended) under Article 34 before the International Preliminary Examining Authority. The description and drawings may only be amended under Article 34 before the International Examining Authority.

Upon entry into the national phase, all parts of the international application may be amended under Article 28 or, where applicable, Article 41.

When?

Within 2 months from the date of transmittal of the international search report or 16 months from the priority date, whichever time limit expires later. It should be noted, however, that the amendments will be considered as having been received on time if they are received by the International Bureau after the expiration of the applicable time limit but before the completion of the technical preparations for international publication (Rule 46.1).

Where not to file the amendments?

The amendments may only be filed with the International Bureau and not with the receiving Office or the International Searching Authority (Rule 46.2).

Where a demand for international preliminary examination has been/is filed, see below.

How?

Either by cancelling one or more entire claims, by adding one or more new claims or by amending the text of one or more of the claims as filed.

A replacement sheet must be submitted for each sheet of the claims which, on account of an amendment or amendments, differs from the sheet originally filed.

All the claims appearing on a replacement sheet must be numbered in Arabic numerals. Where a claim is cancelled, no renumbering of the other claims is required. In all cases where claims are renumbered, they must be renumbered consecutively (Administrative Instructions, Section 205(b)).

The amendments must be made in the language in which the international application is to be published.

What documents must/may accompany the amendments?

Letter (Section 205(b)):

The amendments must be submitted with a letter.

The letter will not be published with the international application and the amended claims. It should not be confused with the "Statement under Article 19(1)" (see below, under "Statement under Article 19(1)").

The letter must be in English or French, at the choice of the applicant. However, if the language of the international application is English, the letter must be in English; if the language of the international application is French, the letter must be in French.

The letter must indicate the differences between the claims as filed and the claims as amended. It must, in particular, indicate, in connection with each claim appearing in the international application (it being understood that identical indications concerning several claims may be grouped), whether

- (i) the claim is unchanged;
- (ii) the claim is cancelled;
- (iii) the claim is new;
- (iv) the claim replaces one or more claims as filed;
- (v) the claim is the result of the division of a claim as filed.

The following examples illustrate the manner in which amendments must be explained in the accompanying letter:

1. [Where originally there were 48 claims and after amendment of some claims there are 51]:
"Claims 1 to 29, 31, 32, 34, 35, 37 to 48 replaced by amended claims bearing the same numbers; claims 30, 33 and 36 unchanged; new claims 49 to 51 added."
2. [Where originally there were 15 claims and after amendment of all claims there are 11]:
"Claims 1 to 15 replaced by amended claims 1 to 11."
3. [Where originally there were 14 claims and the amendments consist in cancelling some claims and in adding new claims]:
"Claims 1 to 6 and 14 unchanged; claims 7 to 13 cancelled; new claims 15, 16 and 17 added." or
"Claims 7 to 13 cancelled; new claims 15, 16 and 17 added; all other claims unchanged."
4. [Where various kinds of amendments are made]:
"Claims 1-10 unchanged; claims 11 to 13, 18 and 19 cancelled; claims 14, 15 and 16 replaced by amended claim 14; claim 17 subdivided into amended claims 15, 16 and 17; new claims 20 and 21 added."

"Statement under article 19(1)" (Rule 46.4)

The amendments may be accompanied by a statement explaining the amendments and indicating any impact that such amendments might have on the description and the drawings (which cannot be amended under Article 19(1)).

The statement will be published with the international application and the amended claims.

It must be in the language in which the international application is to be published.

It must be brief, not exceeding 500 words if in English or if translated into English.

It should not be confused with and does not replace the letter indicating the differences between the claims as filed and as amended. It must be filed on a separate sheet and must be identified as such by a heading, preferably by using the words "Statement under Article 19(1)."

It may not contain any disparaging comments on the international search report or the relevance of citations contained in that report. Reference to citations, relevant to a given claim, contained in the international search report may be made only in connection with an amendment of that claim.

Consequence if a demand for international preliminary examination has already been filed

If, at the time of filing any amendments and any accompanying statement, under Article 19, a demand for international preliminary examination has already been submitted, the applicant must preferably, at the time of filing the amendments (and any statement) with the International Bureau, also file with the International Preliminary Examining Authority a copy of such amendments (and of any statement) and, where required, a translation of such amendments for the procedure before that Authority (see Rules 55.3(a) and 62.2, first sentence). For further information, see the Notes to the demand form (PCT/IPEA/401).

Consequence with regard to translation of the international application for entry into the national phase

The applicant's attention is drawn to the fact that, upon entry into the national phase, a translation of the claims as amended under Article 19 may have to be furnished to the designated/elected Offices, instead of, or in addition to, the translation of the claims as filed.

For further details on the requirements of each designated/elected Office, see Volume II of the PCT Applicant's Guide.

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 98,664-A	FOR FURTHER ACTION see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
International application No. PCT/US 99/ 26788	International filing date (day/month/year) 10/11/1999	(Earliest) Priority Date (day/month/year) 10/11/1998
Applicant DIGITAL GENE TECHNOLOGIES, INC ET AL		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 9 sheets.

☐ It is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

a. With regard to the language, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

b. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☒ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☐ furnished subsequently to this Authority in written form.

☒ furnished subsequently to this Authority in computer readable form.

☒ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☒ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☒ Certain claims were found unsearchable (See Box I).

3. ☒ Unity of invention is lacking (see Box II).

4. With regard to the title,

☐ the text is approved as submitted by the applicant.

☒ the text has been established by this Authority to read as follows:

NUCLEIC ACIDS AND POLYPEPTIDES HAVING ALTERED EXPRESSION IN THE T84 GUT BARRIER MODEL

5. With regard to the abstract,

☐ the text is approved as submitted by the applicant.

☒ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the drawings to be published with the abstract is Figure No.

☐ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

☒ None of the figures.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 99/26788

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☒ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-32 all partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 29 as far as it concerns an in vivo method is directed to a diagnostic method practised on the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Although claims 27 and 28 are directed to a method of treatment of the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box I.2

It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-32 all partially

A polypeptide IMX4 represented by amino acid sequence ID no.27 and DNA molecules represented by nucleic acid sequences ID no.1 and ID no.11,12 and 13. Host cell. Antibody. Variants thereof. Therapeutic uses.

2. Claims: 1-32 all partially

A polypeptide IMX10 represented by amino acid sequence ID no.28 and DNA molecules represented by nucleic acid sequences ID no.2 and ID no.14. Host cell. Antibody. Variants thereof. Therapeutic uses.

3. Claims: 1-32 all partially

A polypeptide IMX21 represented by amino acid sequence ID no.29 and DNA molecules represented by nucleic acid sequences ID no.3 and ID no.15. Host cell. Antibody. Variants thereof. Therapeutic uses.

4. Claims: 1-32 all partially

A polypeptide IMX28 represented by amino acid sequence ID no.30 and DNA molecules represented by nucleic acid sequences ID no.4 and no.16. Host cell. Antibody. Variants thereof. Therapeutic uses.

5. Claims: 1-32 all partially

A polypeptide IMX32 represented by amino acid sequence ID no.31 and DNA molecules represented by nucleic acid sequences ID no.5, no.17 and no.18. Host cell. Antibody. Variants thereof. Therapeutic uses.

6. Claims: 1-32 all partially

A polypeptide IMX39 represented by amino acid sequence ID no.33 and DNA molecules represented by nucleic acid sequences ID no.20 and no.6. Host cell. Antibody. Variants thereof. Therapeutic uses.

7. Claims: 1-32 all partially

A polypeptide IMX40 represented by amino acid sequence ID no.34 and DNA molecules represented by nucleic acid sequences ID no.7 and ID no.21. Host cell. Antibody. Variants thereof. Therapeutic uses.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

8. Claims: 1-32 all partially

A polypeptide IMX42 represented by amino acid sequence ID no.35 and DNA molecules represented by nucleic acid sequences ID no.8. Host cell. Antibody. Variants thereof. Therapeutic uses.

9. Claims: 1-32 all partially

A polypeptide IMX44 represented by amino acid sequence ID no.37 and DNA molecules represented by nucleic acid sequences ID no.9, no.23 and no.24. Host cell. Antibody. Variants thereof. Therapeutic uses.

10. Claims: 1-32 all partially

A polypeptide IMX56 represented by amino acid sequence ID no.38 and DNA molecules represented by nucleic acid sequences ID no.10 and no.25. Host cell. Antibody. Variants thereof. Therapeutic uses.

Box III TEXT OF THE ABSTRACT (Continuation of item 5 of the first sheet)

The invention is directed to purified and isolated polypeptides, the nucleic acids encoding such polypeptides that have altered expression characteristics in the T84 gut barrier model, processes for production of recombinant forms of such polypeptides, antibodies generated against these polypeptides, fragmented peptides derived from these polypeptides and the uses to study inflammatory bowel diseases (IBD).

INTERNATIONAL SEARCH REPORT

International App. No.
PCT/US 99/26788

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12N15/12 C12Q1/68 A61K38/17 C07K14/47 G01N33/68
C07K16/18 A61P1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. HILLIER ET AL: "washU-Merck EST project 1997. zx87d07.r1 Soares tumor NbHOT Homo sapiens cDNA clone IMAGE:810733 5', mRNA sequence" EMBL DATABASE ENTRY HS1272990, ACCESSION NUMBER AA480830, 24 June 1997 (1997-06-24), XP002136972 & UNPUBLISHED, (97% identity with the entire seq ID no.1) --- -/--	4,8,9, 12,15-17



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

4 May 2000

Date of mailing of the international search report

16. 08. 2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

LE CORNEC N.D.R.

INTERNATIONAL SEARCH REPORT

International App. on No

P. US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>P.N. DUCHATEAU ET AL: "Homo sapiens apolipoprotein L mRNA, complete cds" EMBL DATABASE ENTRY AF019225, ACCESSION NUMBER AF019225, 29 September 1997 (1997-09-29), XP002136973 cited in the application -& P.N. DUCHATEAU ET AL: "Apolipoprotein L, a new human high density lipoprotein apolipoprotein expressed by the pancreas. Identification, cloning, characterization, and plasma distribution" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 272, no. 41, 1997, pages 25576-25582, XP002136974 AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD., US ISSN: 0021-9258</p> <p>---</p>	8-17
X	<p>D. LLOYD: "Human DNA sequence from clone RP1-6802 on chromosome 22 contains the 5' end of the APOL2 gene for apolipoprotein L2, the APOL gene for apolipoprotein L, the MYH9 for nonmuscle type myosin heavy chain 9. ESTs, STSs and GSSs" EMBL DATABASE ENTRY HS6802, ACCESSION NUMBER Z82215, 8 November 1996 (1996-11-08), XP002136975 " 99,1% identity with whole sequence ID no.11. Reverse orientation" abstract</p> <p>---</p>	8-17
A	<p>F. BESANCON ET AL: "Interferon-gamma downregulates CFTR gene expression in epithelial cells" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 267, no. 5, November 1994 (1994-11), pages C1398-C1404, XP000906853 the whole document</p> <p>---</p>	1-32
A	<p>J.L. MADARA ET AL: "Interferon-gamma directly affects barrier function of cultured epithelial monolayers" THE JOURNAL OF CLINICAL INVESTIGATION, vol. 83, no. 2, February 1989 (1989-02), pages 724-727, XP000906849 cited in the application the whole document</p> <p>---</p> <p style="text-align: center;">-/--</p>	1-32

INTERNATIONAL SEARCH REPORT

International App.

on No

US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	K.DHARMASATHAPHORN ET AL: "A human colonic tumor cell line that maintains vectorial electrolyte transport" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 9, no. 2, February 1984 (1984-02), pages G204-G208, XP002136976 cited in the application	
A	--- J.L. MADARA ET AL: "Occluding junction structure-function relationships in a cultured epithelial monolayer" THE JOURNAL OF CELL BIOLOGY, vol. 101, December 1985 (1985-12), pages 2124-2133, XP000906833 cited in the application	
T	--- A. YOUAKIM ET AL: "Interferon-gamma decreases barrier function in T84 cells by reducing ZO-1 levels and disrupting apical actin" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 276, no. 5, May 1999 (1999-05), pages G1279-G1288, XP000906855 -----	

PATENT COOPERATION TREATY

DOCKETED

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

PCT

JUL 07 2000

To:

Zimmerman, Roger P.
McDONNELL BOEHNEN HULBERT
& BERGHOFF
300 South Wacker Drive, Suite 3200
CHICAGO, IL 60606
ETATS-UNIS D'AMERIQUE

NOTIFICATION OF RECEIPT OF DEMAND BY COMPETENT INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

(PCT Rules 59.3(e) and 61.1(b), first sentence
and Administrative Instructions, Section 601(a))

Date of mailing
(day/month/year)

23.06.00

Applicant's or agent's file reference

98,664-A

IMPORTANT NOTIFICATION

International application No.

PCT/US 99/26788

International filing date (day/month/year)

10/11/1999

Priority date (day/month/year)

10/11/1998

Applicant

DIGITAL GENE TECHNOLOGIES, INC ET AL

1. The applicant is hereby notified that this International Preliminary Examining Authority considers the following date as the date of receipt of the demand for international preliminary examination of the international application:

06/06/2000

2. This date of receipt is:

- ☒ the actual date of receipt of the demand by this Authority (Rule 61.1(b)).
- ☐ the actual date of receipt of the demand on behalf of this Authority (Rule 59.3(e)).
- ☐ the date on which this Authority has, in response to the invitation to correct defects in the demand (Form PCT/IPEA/404), received the required corrections.

3. ☐ **ATTENTION:** That date of receipt is **AFTER** the expiration of 19 months from the priority date. Consequently, the election(s) made in the demand does (do) not have the effect of postponing the entry into the national phase until 30 months from the priority date (or later in some Offices) (Article 39(1)). Therefore, the acts for entry into the national phase must be performed within 20 months from the priority date (or later in some Offices) (Article 22). For details, see the *PCT Applicant's Guide*, Volume II.

- ☐ (If applicable) This notification confirms the information given by telephone, facsimile transmission or in person on:

4. Only where paragraph 3 applies, a copy of this notification has been sent to the International Bureau.

Name and mailing address of the IPEA/



European Patent Office
D-80298 Munich
Tel. (+49-89) 2399-0, Tx: 523656 epmu d
Fax: (+49-89) 2399-4465

Authorized officer

ALTMANN E F A

Tel. (+49-89) 2399-2378



PATENT COOPERATION TREATY

DOCKETED

From the:
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

Zimmerman, Roger P.
McDONNELL BOEHNEN HULBERT
& BERGHOFF
300 South Wacker Drive, Suite 3200
CHICAGO, IL 60606
ETATS-UNIS D'AMERIQUE

PCT

NOV 14 2000

DUE DATE: 1/3/01

BY: HEC

WRITTEN OPINION

(PCT Rule 66)

Date of mailing
(day/month/year)

09.11.2000

Applicant's or agent's file reference

98,664-A

REPLY DUE

within 2 month(s)

from the above date of mailing

International application No.

PCT/US99/26788

International filing date (day/month/year)

10/11/1999

Priority date (day/month/year)

10/11/1998

International Patent Classification (IPC) or both national classification and IPC

C12N15/12

Applicant

DIGITAL GENE TECHNOLOGIES, INC ET AL

1. This written opinion is the first drawn up by this International Preliminary Examining Authority.

2. This opinion contains indications relating to the following items:

- I ☒ Basis of the opinion
- II ☐ Priority
- III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain document cited
- VII ☒ Certain defects in the international application
- VIII ☒ Certain observations on the international application

3. The applicant is hereby invited to reply to this opinion.

When? See the time limit indicated above. The applicant may, before the expiration of that time limit, request this Authority to grant an extension, see Rule 66.2(d).

How? By submitting a written reply, accompanied, where appropriate, by amendments, according to Rule 66.3. For the form and the language of the amendments, see Rules 66.8 and 66.9.

Also: For an additional opportunity to submit amendments, see Rule 66.4.
For the examiner's obligation to consider amendments and/or arguments, see Rule 66.4 bis.
For an informal communication with the examiner, see Rule 66.6.

If no reply is filed, the international preliminary examination report will be established on the basis of this opinion.

4. The final date by which the international preliminary examination report must be established according to Rule 69.2 is: 10/03/2001.

Name and mailing address of the international preliminary examining authority:

 European Patent Office
D-80298 Munich
Tel. +49 89 2399 - 0 Tx: 523656 epmu d
Fax: +49 89 2399 - 4465

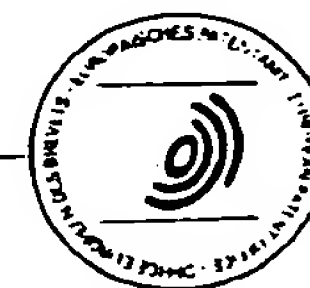
Authorized officer / Examiner

Chavanne, F

Formalities officer (incl. extension of time limits)

Sülberg, A

Telephone No. +49 89 2399 7548



I. Basis of the opinion

1. This opinion has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this opinion as "originally filed".*):

Description, pages:

1-64 as originally filed

Claims, No.:

1-32 as originally filed

Drawings, sheets:

1/23-23/23 as originally filed

2. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
☐ the claims, Nos.:
☐ the drawings, sheets:

3. This opinion has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been and will not be examined in respect of:

- ☐ the entire international application,
☒ claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 4, 6-12, 15-17, 29 (partially),

because:

- ☐ the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (*specify*):

- ☐ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):
- ☒ the claims, or said claims Nos. 1-3, 5, 13, 14, 18-28, 30-32 (completely), 9-12, 15-17 (partially) are so inadequately supported by the description that no meaningful opinion could be formed.
- ☒ no international search report has been established for the said claims Nos. 1-32 (partially).

V. Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Claims	9-12, 15-17
Inventive step (IS)	Claims	4, 6-12, 15-17, 29
Industrial applicability (IA)	Claims	

2. Citations and explanations

see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. As a result of an objection of lack of unity raised by the International Search Authority, the international search report has been restricted to the invention referred to as invention 1 (claims 1-32, partially), namely:
a polypeptide IMX4 represented by amino acid sequence ID No. 27 and DNA molecules represented by nucleic acid sequences ID No. 1 and ID No. 11, 12 and 13, antibodies binding to said polypeptide, host cells expressing said polypeptide, variants thereof, and therapeutic uses thereof.
Consequently, an opinion with regard to novelty, inventive step and industrial applicability can only be partially formulated, and limited to the above mentioned invention 1.
2. An opinion with regard to novelty, inventive step and industrial applicability cannot be formulated for the claims relating to a polypeptide comprising the amino acid sequence of SEQ ID No. 27, or a polypeptide encoded by a polynucleotide of SEQ ID No. 1, for the following reasons:

According to the present application, the polypeptide having the amino acid sequence of SEQ ID No. 27 is the translational product of the nucleotide sequence of isolated DNA molecule IMX4 of SEQ ID No. 1 (see e.g. page 59, example 2). However, in view of the nucleotide sequence of SEQ ID No. 1, it appears that said DNA molecule cannot encode said polypeptide:

- A. The nucleotide sequence of SEQ ID No. 1 contains only one putative methionine initiation codon (ATG) at position 29. The corresponding putative open reading frame (ORF) stops after three amino acids (KAV) with a stop codon (TAG) at position 41. This is too short to corresponds to an ORF leading to a translated product. Thus, there are at the nucleotide level no indications that the nucleotide sequence of SEQ ID No. 1 encodes for a polypeptide.
- B. The comparison of the amino acid sequence of SEQ ID No. 27 with the nucleotide sequence of SEQ ID No. 1 shows that the codon supposed to

correspond to the methionine of SEQ ID No. 27 actually corresponds to a stop codon (TAG). Thus, the polypeptide of SEQ ID No. 27 cannot be translated from IMX4 as asserted in the present application (see e.g. page 59, line 15).

- C. The alignment of the nucleotide sequence of SEQ ID No. 1 with the nucleotide sequence of the soares ovary tumour HbHOT Homo sapiens cDNA clone (D1) shows that the nucleotide sequence of SEQ ID No. 1 corresponds to a 3' untranslated region. In fact, the nucleotide sequence of SEQ ID No. 1 shows over 96% identity with the 3' untranslated region of said human cDNA clone. This alignment shows that the last five A nucleotides of SEQ ID No. 1 are part of the polyA tail. A polyadenylation signal motif can also be found at position 114. Moreover, because of the high mutation rate of non-coding sequences, 3' untranslated regions are gene specific. Thus, it can be totally excluded that a 3' untranslated region would show such a high identity with a coding region.
- Therefore, the nucleotide sequence of SEQ ID No. 1, which is supposed to encode the polypeptide having the amino acid sequence of SEQ ID No. 27, actually corresponds to a 3' untranslated region.

Since the polypeptide of SEQ ID No. 27 does not exist, and since the nucleotide sequence of SEQ ID No. 1 does not encode any polypeptide, an opinion with regard to novelty, inventive step and industrial applicability on the claims referring to said polypeptides, and related claims cannot be formulated (claims 1-3, 5, 13, 14, 18-28 and 30-32, completely; 9-12 and 15-17, partially).

V. Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

D1: EMBL Database entry HS1272990
Accession Number AA480830

D2: EMBL Database entry HS6802
Accession Number Z82215.1

2. D1 describes the nucleotide sequence of a cDNA clone comprising a sequence showing over 96% identity with SEQ ID No. 1.
D2 describes a nucleotide sequence showing over 97% identity with the nucleotide sequence of SEQ ID No. 1.
Thus, in view of D1 or D2, the subject-matter of claims 9-12 and 15-17 is not novel. Therefore, claims 9-12 and 15-17 do not meet the requirements of Article 33(2) PCT.
3. The nucleotide sequence of SEQ ID No. 1 described in the present application corresponds to the 3' untranslated region of an unknown gene. The present application shows that the transcription of said gene is increased when cells are treated with interferon- γ (example 1). However, this does not give any evidence on the specific function of the product of said gene. The function of the product of the gene corresponding to the 3' untranslated region of nucleotide sequence of SEQ ID No.1 being unknown, the isolation of said nucleotide does not solve any problem. Since there is no problem to be solved, there cannot be any inventive step involved. Thus, the subject-matter of claims 4, 6-12, 15-17 and 29 is not inventive. Therefore, claims 4, 6-12, 15-17 and 29 do not meet the requirements of Article 33(3) PCT.

VII. Certain defects in the international application

1. Point (d) of claim 9 is lacking.
2. The specification mentions that the sequences of SEQ ID No. 11 and 12 include the sequence of SEQ ID No. 1 (page 59, lines 7-10). It appears that this is not the case.
3. The specification does not define the SEQ ID No. 18, 19 and 21-26 although they are mentioned in the claims.

4. Claim 16 refers to a method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 9. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject. Both claims do not mention how said methods are to be performed.

VIII. Certain observations on the international application

1. The formulation "...comprises..." in claims 4 and 8-12 does not clearly define the scope of the claim. Thus, this expression should be replaced with "consists of" (Article 6 PCT).
2. Claims 9 and 12 lack clarity due to the formulation "hybridisable to". This formulation is vague and imprecise, it does not provide any technical information about the hybridisation conditions, and as a consequence is not adapted to clearly define the scope of the claim. In fact, any nucleic acid molecule may hybridise to the nucleic acid molecule of SEQ ID No. 1, when the hybridisation conditions are adapted to their degree of identity (Article 6 PCT).
3. Claim 1 mentions that the DNA molecule of SEQ ID No. 1 has altered expression in a T84 model of gut barrier function. However, the specification does not provide any evidence for said altered expression. The present application only shows that treatment of the T84 cells with interferon- γ induces expression of the DNA molecule of SEQ ID No. 1. Thus, claim 1 is not based on the description (Article 6-support PCT).
4. Claim 29 refers to a method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject by determining the presence or absence of a polynucleotide. Apart from vague declarations, the present application does not provide any evidence showing that the irritable bowel disease may be evidenced by determining the presence or absence of the polynucleotide of SEQ ID No. 1. Thus, claim 29 is not based on the description (Article 6-support PCT). Moreover, since a polynucleotide is always either present or absent, the man skilled in the art, faced with the problem of establishing such a diagnosis would not be able to perform it with said method (Article 5 PCT).

**IN THE INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY
EUROPEAN PATENT OFFICE**

International application No. PCT/US99/26788	International filing date (day/month/year) 10 November 1999 (10.11.99)	(Earliest) Priority date (day/month/year) 10 November 1998 (10.11.98)
Title of invention NOVEL DNAS AND POLYPEPTIDES		
Applicant(s) DIGITAL GENE TECHNOLOGIES, INC.		

TRANSMITTAL LETTER

IPEA
European Patent Office
D-80298 Munich
Germany

Attn: Examiner F. Chavanne

Sir:

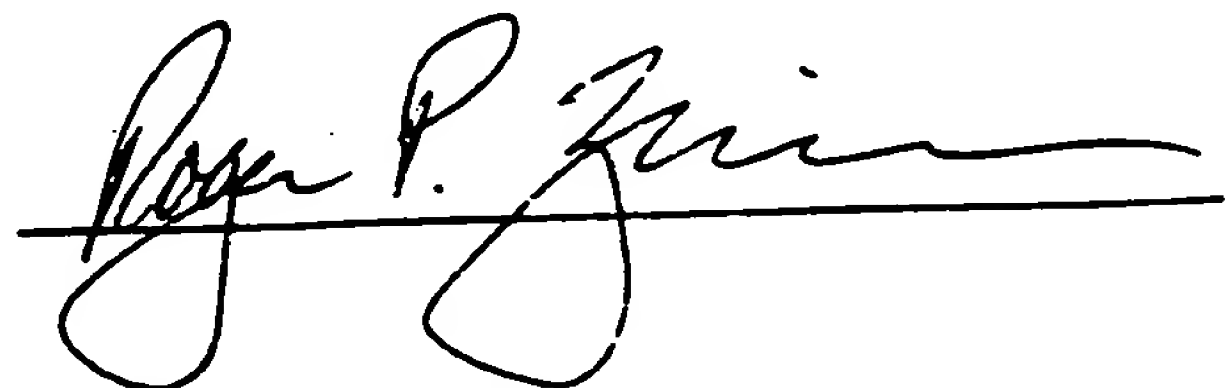
In regard to the above identified application:

1. We are transmitting herewith by DHL Courier the attached:
 - A. Response to the Written Opinion with Exhibits A-C;
 - B. Substitute Specification and Claims pgs. 1-74;
2. We are transmitting herewith by facsimile:
3.
 - A. Response to Written Opinion with Exhibits A-C
3. With respect to additional fees, no additional fee is required.

By :


Roger P. Zimmerman
U.S. Patent Office Reg. No. 38,670

CERTIFICATE OF MAILING BY "FACSIMILE WITH CONFIRMATION BY DHL COURIER": The undersigned hereby certifies that this Transmittal Letter and the paper, as described in paragraph 2 hereinabove, are being transmitted by Facsimile to the European Patent Office, Attn: F. Chavanne, Fax: 011-49-89-2399-4465, and deposited with the DHL Courier as described by paragraph 1 herein above with sufficient postage in an envelope addressed to: European Patent Office, Attn: F. Chavanne, D-80298 Munich, Germany, on this 7th day of February, 2001. DHL No. 8377549902.



INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 99/26788

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12N15/12 C12Q1/68 A61K38/17 C07K14/47 G01N33/68
C07K16/18 A61P1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. HILLIER ET AL: "washU-Merck EST project 1997. zx87d07.r1 Soares tumor NbHOT Homo sapiens cDNA clone IMAGE:810733 5', mRNA sequence" EMBL DATABASE ENTRY HS1272990, ACCESSION NUMBER AA480830, 24 June 1997 (1997-06-24), XP002136972 & UNPUBLISHED, (97% identity with the entire seq ID no.1) --- -/--	4,8,9, 12,15-17

☒ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

4 May 2000

Date of mailing of the international search report

16. 08. 2000

Name and mailing address of the ISA

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Authorized officer

LE CORNEC N.D.R.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>P.N. DUCHATEAU ET AL: "Homo sapiens apolipoprotein L mRNA, complete cds" EMBL DATABASE ENTRY AF019225, ACCESSION NUMBER AF019225, 29 September 1997 (1997-09-29), XP002136973 cited in the application -& P.N. DUCHATEAU ET AL: "Apolipoprotein L, a new human high density lipoprotein apolipoprotein expressed by the pancreas. Identification, cloning, characterization, and plasma distribution" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 272, no. 41, 1997, pages 25576-25582, XP002136974 AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD., US ISSN: 0021-9258</p> <p>---</p>	8-17
X	<p>D. LLOYD: "Human DNA sequence from clone RP1-6802 on chromosome 22 contains the 5' end of the APOL2 gene for apolipoprotein L2, the APOL gene for apolipoprotein L, the MYH9 for nonmuscle type myosin heavy chain 9. ESTs, STSs and GSSs" EMBL DATABASE ENTRY HS6802, ACCESSION NUMBER Z82215, 8 November 1996 (1996-11-08), XP002136975 " 99,1% identity with whole sequence ID no.11. Reverse orientation" abstract</p> <p>---</p>	8-17
A	<p>F. BESANÇON ET AL: "Interferon-gamma downregulates CFTR gene expression in epithelial cells" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 267, no. 5, November 1994 (1994-11), pages C1398-C1404, XP000906853 the whole document</p> <p>---</p>	1-32
A	<p>J.L. MADARA ET AL: "Interferon-gamma directly affects barrier function of cultured epithelial monolayers" THE JOURNAL OF CLINICAL INVESTIGATION, vol. 83, no. 2, February 1989 (1989-02), pages 724-727, XP000906849 cited in the application the whole document</p> <p>---</p>	1-32
	<p>---</p> <p>-/--</p>	

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/26788

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	K.DHARMASATHAPHORN ET AL: "A human colonic tumor cell line that maintains vectorial electrolyte transport" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 9, no. 2, February 1984 (1984-02), pages G204-G208, XP002136976 cited in the application	
A	--- J.L. MADARA ET AL: "Occluding junction structure-function relationships in a cultured epithelial monolayer" THE JOURNAL OF CELL BIOLOGY, vol. 101, December 1985 (1985-12), pages 2124-2133, XP000906833 cited in the application	
T	--- A. YOUAKIM ET AL: "Interferon-gamma decreases barrier function in T84 cells by reducing ZO-1 levels and disrupting apical actin" AMERICAN JOURNAL OF PHYSIOLOGY, vol. 276, no. 5, May 1999 (1999-05), pages G1279-G1288, XP000906855 -----	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 99/26788

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☒ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-32 all partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 29 as far as it concerns an in vivo method is directed to a diagnostic method practised on the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Although claims 27 and 28 are directed to a method of treatment of the human/animal body (rule 39.1 IV PCT), the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box I.2

It has to be noted that sequences ID no.19, no.22, no.26, no.32 and no.36 do not exist in the originally filed sequence listing on which was based the search. There are in total 33 sequences and not 38. But the originally numbering of the sequences has been kept.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-32 all partially

A polypeptide IMX4 represented by amino acid sequence ID no.27 and DNA molecules represented by nucleic acid sequences ID no.1 and ID no.11,12 and 13. Host cell. Antibody. Variants thereof. Therapeutic uses.

2. Claims: 1-32 all partially

A polypeptide IMX10 represented by amino acid sequence ID no.28 and DNA molecules represented by nucleic acid sequences ID no.2 and ID no.14. Host cell. Antibody. Variants thereof. Therapeutic uses.

3. Claims: 1-32 all partially

A polypeptide IMX21 represented by amino acid sequence ID no.29 and DNA molecules represented by nucleic acid sequences ID no.3 and ID no.15. Host cell. Antibody. Variants thereof. Therapeutic uses.

4. Claims: 1-32 all partially

A polypeptide IMX28 represented by amino acid sequence ID no.30 and DNA molecules represented by nucleic acid sequences ID no.4 and no.16. Host cell. Antibody. Variants thereof. Therapeutic uses.

5. Claims: 1-32 all partially

A polypeptide IMX32 represented by amino acid sequence ID no.31 and DNA molecules represented by nucleic acid sequences ID no.5, no.17 and no.18. Host cell. Antibody. Variants thereof. Therapeutic uses.

6. Claims: 1-32 all partially

A polypeptide IMX39 represented by amino acid sequence ID no.33 and DNA molecules represented by nucleic acid sequences ID no.20 and no.6. Host cell. Antibody. Variants thereof. Therapeutic uses.

7. Claims: 1-32 all partially

A polypeptide IMX40 represented by amino acid sequence ID no.34 and DNA molecules represented by nucleic acid sequences ID no.7 and ID no.21. Host cell. Antibody. Variants thereof. Therapeutic uses.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

8. Claims: 1-32 all partially

A polypeptide IMX42 represented by amino acid sequence ID no.35 and DNA molecules represented by nucleic acid sequences ID no.8. Host cell. Antibody. Variants thereof. Therapeutic uses.

9. Claims: 1-32 all partially

A polypeptide IMX44 represented by amino acid sequence ID no.37 and DNA molecules represented by nucleic acid sequences ID no.9, no.23 and no.24. Host cell. Antibody. Variants thereof. Therapeutic uses.

10. Claims: 1-32 all partially

A polypeptide IMX56 represented by amino acid sequence ID no.38 and DNA molecules represented by nucleic acid sequences ID no.10 and no.25. Host cell. Antibody. Variants thereof. Therapeutic uses.

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C12N 15/12, C12Q 1/68, A61K 38/17, C07K 14/47, G01N 33/68, C07K 16/18, A61P 1/00	A2	(11) International Publication Number: WO 00/28033 (43) International Publication Date: 18 May 2000 (18.05.00)
(21) International Application Number: PCT/US99/26788 (22) International Filing Date: 10 November 1999 (10.11.99) (30) Priority Data: 60/107,821 10 November 1998 (10.11.98) US (71) Applicant (for all designated States except US): DIGITAL GENE TECHNOLOGIES, INC. [US/US]; Suite 110, 11149 North Torrey Pines Road, La Jolla, CA 92037 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): BAUM, Peter, Robert [US/US]; 6027 34th Avenue NE, Seattle, WA 98115 (US). DUBOSE, Robert, Finley [US/US]; 2981 142nd Place NE, Bellevue, WA 98007 (US). SIMS, John, E. [US/US]; 4207 43rd Avenue NE, Seattle, WA 98105 (US). YOUAKIM, Adel [CA/US]; 7304 175th Street SW, Edmonds, WA 98026 (US). HASEL, Karl, W. [US/US]; 247 South Nardo Avenue, Solana Beach, CA 92075 (US). HILBUSH, Brian, S. [US/US]; 13058 Caminito Angelico, San Diego, CA 92130 (US).		(74) Agent: ZIMMERMAN, Roger, P.; McDonnell Boehnen Hul- bert & Berghoff, 32nd floor, 300 South Wacker Drive, Chicago, IL 60606 (US). (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: NOVEL DNAs AND POLYPEPTIDES (57) Abstract The invention is directed to purified and isolated novel polypeptides, the nucleic acids encoding such polypeptides, processes for production of recombinant forms of such polypeptides, antibodies generated against these polypeptides, fragmented peptides derived from these polypeptides, and the uses of the above.		

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C12N 15/12, C12Q 1/68, A61K 38/17, C07K 14/47, G01N 33/68, C07K 16/18, A61P 1/00	A2	(11) International Publication Number: WO 00/28033 (43) International Publication Date: 18 May 2000 (18.05.00)
(21) International Application Number: PCT/US99/26788 (22) International Filing Date: 10 November 1999 (10.11.99) (30) Priority Data: 60/107,821 10 November 1998 (10.11.98) US (71) Applicant (for all designated States except US): DIGITAL GENE TECHNOLOGIES, INC. [US/US]; Suite 110, 11149 North Torrey Pines Road, La Jolla, CA 92037 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): BAUM, Peter, Robert [US/US]; 6027 34th Avenue NE, Seattle, WA 98115 (US). DUBOSE, Robert, Finley [US/US]; 2981 142nd Place NE, Bellevue, WA 98007 (US). SIMS, John, E. [US/US]; 4207 43rd Avenue NE, Seattle, WA 98105 (US). YOUAKIM, Adel [CA/US]; 7304 175th Street SW, Edmonds, WA 98026 (US). HASEL, Karl, W. [US/US]; 247 South Nardo Avenue, Solana Beach, CA 92075 (US). HILBUSH, Brian, S. [US/US]; 13058 Caminito Angelico, San Diego, CA 92130 (US).		(74) Agent: ZIMMERMAN, Roger, P.; McDonnell Boehnen Hulbert & Berghoff, 32nd floor, 300 South Wacker Drive, Chicago, IL 60606 (US). (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: NOVEL DNAs AND POLYPEPTIDES (57) Abstract The invention is directed to purified and isolated novel polypeptides, the nucleic acids encoding such polypeptides, processes for production of recombinant forms of such polypeptides, antibodies generated against these polypeptides, fragmented peptides derived from these polypeptides, and the uses of the above.		

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DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

NOVEL DNAs AND POLYPEPTIDES

Reference to Related Application

This application claims the benefit of U.S. Provisional Application S.N. 60/107,821, filed November 10, 1998, which is incorporated herein by reference.

BACKGROUND OF THE INVENTION

Field of the Invention

The invention is directed to purified and isolated novel polypeptide molecules and fragments thereof, the nucleic acid molecules encoding such polypeptides, processes for production of recombinant forms of such polypeptide molecules, antibodies generated against such polypeptide molecules, fragmented peptides derived from these polypeptide molecules, and uses thereof. In particular, the invention is directed to the use of the nucleic acid and polypeptide molecules in the study of inflammatory bowel diseases (IBD)

Description of Related Art

Damage to the intestinal epithelial barrier is a hallmark of inflammatory bowel diseases (IBD). Examples of inflammatory bowel diseases include ileitis, Crohn's disease (CD), which can affect the whole digestive tract from mouth to anus, and ulcerative colitis (UC), which affects only the large intestine. Studying the factors that influence the integrity of the epithelial barrier *in vivo* is a difficult task for a number of reasons that include the complexity of the tissue itself (there are numerous cell types in the gut including epithelial, stromal, endocrine, neuronal and hematopoietic) and the technical problems associated with tissue manipulation in animals or in isolated organs. As a result of these issues, a number of *in vitro* models of epithelial barrier function have been developed over the years. The best characterized of these models is the T84 intestinal epithelial barrier system (Dharmasathaphorn et al., *Am. J. Physiol.*, 246:G204-G208, 1984 and Madara et al., *J. Cell Biol.*, 101:2124-2133, 1985). T84 cells, while derived from a human colonic adenocarcinoma, have retained many of the properties associated with normal colonic crypt cells. T84 cells form polarized monolayers that exhibit high electrical resistance and vectorial fluid and chloride secretion reminiscent of colonic crypts *in vivo*. These properties are directly dependent on a complex of proteins referred to as tight junctions and T84 cells express many of the known members of this complex (Youakim et al., *Submitted for*

publication, 1998). These cells also respond to proinflammatory cytokines, such as interferon-gamma (INF-gamma), by decreasing barrier function (Youakim et al., *Submitted for publication*, 1998; Madara et al., *J. Clin. Invest.*, 83:724-727, 1989; and Adams et al., *J. Immunol.*, 150:2356-2363, 1993) and by up-regulating MHC Class II molecules and antigen presenting activity (Hershberg et al., *J. Clin. Invest.*, 102:792-803, 1998). This model system is used to examine how the epithelial barrier is regulated by various agents such as interferon-gamma and cells of the immune system (e.g., neutrophils). The T84 model assists in the elucidation of the mechanism of barrier breakdown and recovery in response to these agents and in the identification of proteins (and genes) that may prevent barrier breakdown or stimulate barrier recovery.

Given the important function of the epithelial gut barrier and despite the growing body of knowledge, there is a need in the art for the discovery, the identification, and the elucidation of the roles of new proteins involved in gut barrier function and IBD.

The identification of the primary structure, or sequence, of an unknown protein is the culmination of an arduous process of experimentation. In order to identify an unknown protein, the investigator can rely upon a comparison of the unknown protein to known peptides using a variety of techniques known to those skilled in the art. For instance, proteins are routinely analyzed using techniques such as electrophoresis, sedimentation, chromatography, sequencing and mass spectrometry.

In particular, comparison of an unknown protein to polypeptides of known molecular weight allows a determination of the apparent molecular weight of the unknown protein (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)). Protein molecular weight standards are commercially available to assist in the estimation of molecular weights of unknown protein (New England Biolabs Inc. Catalog:130-131, 1995; J. L. Hartley, U.S. Patent No. 5,449,758). However, the molecular weight standards may not correspond closely enough in size to the unknown protein to allow an accurate estimation of apparent molecular weight. The difficulty in estimation of molecular weight is compounded in the case of proteins that are subjected to fragmentation by chemical or enzymatic means, modified by post-translational modification or processing, and/or associated with other proteins in non-covalent complexes.

In addition, the unique nature of the composition of a protein with regard to its specific amino acid constituents results in unique positioning of cleavage sites within the protein. Specific fragmentation of a protein by chemical or enzymatic cleavage results in a unique "peptide fingerprint" (D. W. Cleveland et al., *J. Biol. Chem.* 252:1102-1106, 1977; M. Brown et

al., *J. Gen. Virol.* 50:309-316, 1980). Consequently, cleavage at specific sites results in reproducible fragmentation of a given protein into peptides of precise molecular weights. Furthermore, these peptides possess unique charge characteristics that determine the isoelectric pH of the peptide. These unique characteristics can be exploited using a variety of electrophoretic and other techniques (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)).

Fragmentation of proteins is further employed for amino acid composition analysis and protein sequencing (P. Matsudaira, *J. Biol. Chem.* 262:10035-10038, 1987; C. Eckerskorn et al., *Electrophoresis* 1988, 9:830-838, 1988), particularly the production of fragments from proteins with a "blocked" N-terminus. In addition, fragmented proteins can be used for immunization, for affinity selection (R. A. Brown, U.S. Patent No. 5,151,412), for determination of modification sites (e.g. phosphorylation), for generation of active biological compounds (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 300-301 (Prentice Hall, 6d ed. 1991)), and for differentiation of homologous proteins (M. Brown et al., *J. Gen. Virol.* 50:309-316, 1980).

In addition, when a peptide fingerprint of an unknown protein is obtained, it can be compared to a database of known proteins to assist in the identification of the unknown protein using mass spectrometry (W.J. Henzel et al., *Proc. Natl. Acad. Sci. USA* 90:5011-5015, 1993; D. Fenyo et al., *Electrophoresis* 19:998-1005, 1998). A variety of computer software programs to facilitate these comparisons are accessible via the Internet, such as Protein Prospector (Internet site: prospector.uscf.edu), MultiIdent (Internet site: www.expasy.ch/sprot/multiident.html), PeptideSearch (Internet site: www.mann.embl-heidelberg.de/deSearch/FR_PeptideSearchForm.html), and ProFound (Internet site: www.chait-sgi.rockefeller.edu/cgi-bin/prot-id-frag.html). These programs allow the user to specify the cleavage agent and the molecular weights of the fragmented peptides within a designated tolerance. The programs compare these molecular weights to protein molecular weight information stored in databases to assist in determining the identity of the unknown protein. Accurate information concerning the number of fragmented peptides and the precise molecular weight of those peptides is required for accurate identification. Therefore, increasing the accuracy in determining the number of fragmented peptides and their molecular weight should result in enhanced likelihood of success in the identification of unknown proteins.

In addition, peptide digests of unknown proteins can be sequenced using tandem mass spectrometry (MS/MS), and the resulting sequence searched against databases (J.K. Eng, et al., *J. Am. Soc. Mass Spec.* 5:976-989 (1994); M. Mann and M. Wilm, *Anal. Chem.* 66:4390-4399

(1994); J.A. Taylor and R.S. Johnson, *Rapid Comm. Mass Spec.* 11:1067-1075 (1997)).

Searching programs that can be used in this process exist on the Internet, such as Lutefisk 97 (Internet site: www.lsbcc.com:70/Lutefisk97.html), and the Protein Prospector, Peptide Search and ProFound programs described above. Therefore, adding the sequence of a gene and its
5 predicted protein sequence and peptide fragments to a sequence database can aid in the identification of unknown proteins using tandem mass spectrometry.

Thus, there also exists a need in the art for polypeptides suitable for use in peptide fragmentation studies, preferably, polypeptides that have altered expression in irritable bowel diseases, for use in molecular weight measurements, and for use in protein sequencing using
10 tandem mass spectrometry.

SUMMARY OF THE INVENTION

Using the T84 model, it was determined that certain polypeptides have altered (up-regulated or down-regulated) expression patterns in response to INF-gamma. Such molecules
15 may have a role in gut barrier function and IBD and may be useful as potential therapeutic agents in the treatment of IBD and other gut pathologies. The invention aids in fulfilling these needs in the art by providing isolated nucleic acids and polypeptides encoded by these nucleic acids that have altered expression characteristics in the T84 gut barrier model. Particular
embodiments of the invention are directed to isolated nucleic acid molecules comprising the
20 DNA sequences of SEQ ID NOs:1-26 and isolated nucleic acid molecules encoding the amino acid sequences of SEQ ID NOs:27-38, as well as nucleic acid molecules complementary to these sequences. Both single-stranded and double-stranded RNA and DNA nucleic acid molecules are encompassed by the invention, as well as nucleic acid molecules that hybridize to a denatured,
double-stranded DNA comprising all or a portion of SEQ ID NOs:1-26 and/or the DNA that
25 encodes the amino acid sequences of SEQ ID NOs:27-38. Also encompassed are isolated nucleic acid molecules that are derived by *in vitro* mutagenesis from nucleic acid molecules comprising sequences of SEQ ID NOs:1-26, that are degenerate from nucleic acid molecules comprising sequences of SEQ ID NOs:1-26, and that are allelic variants of DNA of the
invention. The invention also encompasses recombinant vectors that direct the expression of
30 these nucleic acid molecules and host cells transformed or transfected with these vectors.

In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%, more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO:1-26 or a polynucleotide which is hybridizable to SEQ ID NO:1-26;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 1-26 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO:1-26;

(c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 1-26 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO:1-26;

(e) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 1-26, having biological activity;

(f) a polynucleotide which is a variant of SEQ ID NO:1-26;

(g) a polynucleotide which is an allelic variant of SEQ ID NO:1-26;

(h) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 1-26;

(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%, more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO:1-10 or a polynucleotide which is hybridizable to SEQ ID NO:1-10;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 1-10 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO:1-10;

(c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 1-10 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO:1-10;

(e) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 1-10, having biological activity;

(f) a polynucleotide which is a variant of SEQ ID NO:1-10;

(g) a polynucleotide which is an allelic variant of SEQ ID NO:1-10;

(h) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 1-10;

(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%, more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO:11-26 or a polynucleotide which is hybridizable to SEQ ID NO:11-26;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 11-26 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO:11-26;

(c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 11-26 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO:11-26;

(e) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 11-26, having biological activity;

(f) a polynucleotide which is a variant of SEQ ID NO:11-26;

(g) a polynucleotide which is an allelic variant of SEQ ID NO:11-26;

(h) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 11-26;

(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

In one embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a secreted protein. In preferred embodiments, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a polypeptide chosen from the group consisting of:

(a) a polypeptide having the polypeptide sequence identified as a translation of SEQ ID NO: 1-26;

(b) a polypeptide having the polypeptide sequence of SEQ ID NO: 27-38; and

(c) a polypeptide encoded by the cDNA which is hybridizable to SEQ ID NO:1-26. Typically, the isolated nucleic acid molecule comprises the entire nucleotide sequence of SEQ ID NO:1-26 or a cDNA sequence which is hybridizable to SEQ ID NO:1-26. Typically the isolated nucleic acid molecule comprises sequential nucleotide deletions from portions of the nucleotide sequence encoding either the C-terminus or the N-terminus of the polypeptide.

In other aspects, the invention provides a recombinant vector comprising the isolated nucleic acid molecule and a method of making a recombinant host cell comprising the isolated nucleic acid molecule and the recombinant host cell produced by such a method.

In other preferred embodiments, the invention provides an isolated polypeptide having an amino acid sequence at least 80%, preferably at least 85%, more preferably at least 90% identical to the sequence of a polypeptide selected from the group consisting of:

(a) a polypeptide fragment of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(b) a polypeptide having the sequence of SEQ ID NO: 27-38;

(c) a polypeptide domain of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(d) a polypeptide epitope of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(e) a secreted form of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(f) a full length protein of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(g) a variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(h) an allelic variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

and

(i) a species homologue of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26.

In some embodiments, the full length polypeptide comprises sequential amino acid deletions from the C-terminus. In other embodiments, the mature polypeptide comprises sequential amino acid deletions from the C-terminus. Alternatively, the full length polypeptide can comprise sequential amino acid deletions from the N-terminus or the mature polypeptide can comprise sequential amino acid deletions from the N-terminus.

In other aspects, the invention provides an isolated antibody that binds specifically to the isolated polypeptide, a recombinant host cell that expresses the isolated polypeptide, and a method of making an isolated polypeptide comprising culturing the recombinant host cell under conditions such that said polypeptide is expressed; and recovering said polypeptide.

5

The invention in another embodiment is a method for preventing, treating, or ameliorating irritable bowel disorders, comprising administering to a mammalian subject a therapeutically effective amount of the isolated polypeptide or the isolated nucleic acid molecule. In another embodiment, the invention provides a method of diagnosing an irritable
10 bowel disease or a susceptibility to irritable bowel disease in a subject comprising: determining the presence or absence of a mutation in the isolated nucleic acid molecule, and diagnosing an irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or absence of said mutation. Alternatively, irritable bowel disease or a susceptibility to irritable
15 bowel disease in a subject can be diagnosed by a method comprising: determining the presence or amount of expression of the polypeptide in a biological sample; and diagnosing irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or amount of expression of the polypeptide.

Typically, a binding partner of the polypeptide is identified by contacting the polypeptide with a binding partner; and determining whether the binding partner affects a
20 physical property or an activity of the polypeptide. Typically activity in a biological assay of a secreted polypeptide is identified by expressing the polynucleotide of SEQ ID NO:1-26 in a cell; isolating the supernatant; detecting an activity in a biological assay; and identifying the polypeptide in the supernatant having the activity.

25 In addition, the invention encompasses methods of using the nucleic acids noted above to identify nucleic acids encoding proteins homologous to SEQ ID NOs:27-38; to identify human chromosomes that contain the nucleotide sequences of the invention; to map genes near the nucleotide sequences of the invention on human chromosomes; and to identify genes associated with certain diseases, syndromes, or other human conditions associated with human
30 chromosomes containing sequences of the invention.

For example, four of the nucleotide sequences of the invention, IMX4 and IMX56, IMX21, and IMX44 are located on chromosomes 22, 22, 7, and 19, respectively (IMX4 and IMX56 are both located on chromosome 22). Thus, the above-named nucleotide sequences (IMX4, IMX56, IMX21, and IMX44) can be used to identify human chromosome numbers 22,

7, and 19; to map genes on human chromosome numbers 22, 7, and 19; and to identify genes associated with certain diseases, syndromes, or other human conditions associated with human chromosome numbers 22, 7, and 19.

5 The invention also encompasses the use of sense or antisense oligonucleotides from the nucleic acids of SEQ ID NOs:1-26 to inhibit the expression of the polynucleotides encoded by the nucleotide sequences of the invention.

10 The invention also encompasses isolated polypeptides and fragments thereof encoded by these nucleic acid molecules including soluble polypeptide portions of SEQ ID NOs:27-38. The invention further encompasses methods for the production of these polypeptides, including culturing a host cell under conditions promoting expression and recovering the polypeptide from the culture medium if it is secreted or from cultured cells if it is not secreted. Especially, the expression of these polypeptides in bacteria, yeast, plant, insect, and animal cells is encompassed by the invention.

15 In addition, the invention includes assays utilizing these polypeptides, to screen for potential inhibitors of activity associated with polypeptide counter-structure molecules, and methods of using these polypeptides as therapeutic agents for the treatment of diseases mediated by polypeptide counter-structure molecules. Further, methods of using these polypeptides in the design of inhibitors thereof are also an aspect of the invention.

20 The invention further includes a method for using these polypeptides as molecular weight markers that allow the estimation of the molecular weight of a protein or a fragmented protein, as well as a method for the visualization of the molecular weight markers of the invention thereof using electrophoresis. The invention further encompasses methods for using the polypeptides of the invention as markers for determining the isoelectric point of an unknown protein, as well as controls for establishing the extent of fragmentation of a protein.

25 Further encompassed by this invention are kits to aid in these determinations.

Further encompassed by this invention is the use of the IMX nucleic acid sequences, predicted amino acid sequences of the polypeptide or fragments thereof, or a combination of the predicted amino acid sequences of the polypeptides and fragments thereof for use in searching an electronic database to aid in the identification of sample nucleic acids and/or proteins.

30 The invention also encompasses IMX polypeptides and the use of these polypeptides as research reagents to further study gut epithelial barrier function and regulation and therapeutic reagents to treat inflammatory bowel disease and other gut pathologies.

Isolated polyclonal or monoclonal antibodies that bind to these polypeptides are also encompassed by the invention, in addition the use of these antibodies to aid in purifying IMX polypeptides.

5

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 presents the nucleotide sequence of IMX 4 (SEQ ID NO: 1).

Figure 2 presents the nucleotide sequence of IMX 10 (SEQ ID NO: 14).

Figure 3 presents the nucleotide sequence of IMX 21 (SEQ ID NO: 15).

Figure 4 presents the nucleotide sequence of IMX 28 (SEQ ID NO: 16).

10

Figure 5 presents the nucleotide sequence of IMX 32 (SEQ ID NO: 17).

Figure 6 presents the nucleotide sequence of IMX 39 (SEQ ID NO: 20).

Figure 7 presents the nucleotide sequence of IMX 40 (SEQ ID NO: 7).

Figure 8 presents the nucleotide sequence of IMX 42 (SEQ ID NO: 8).

Figure 9 presents the nucleotide sequence of IMX 44 (SEQ ID NO: 23).

15

Figure 10 presents the nucleotide sequence of IMX 56 (SEQ ID NO: 10).

Figure 11 presents the amino acid sequence of IMX 4 (SEQ ID NO: 27).

Figure 12 presents the amino acid sequence of IMX 10 (SEQ ID NO: 28).

Figure 13 presents the amino acid sequence of IMX 21 (SEQ ID NO: 29).

Figure 14 presents the amino acid sequence of IMX 28 (SEQ ID NO: 30).

20

Figure 15 presents the amino acid sequence of IMX 32 (SEQ ID NO: 32).

Figure 16 presents the amino acid sequence of IMX 39 (SEQ ID NO: 33).

Figure 17 presents the amino acid sequence of IMX 40 (SEQ ID NO: 34).

Figure 18 presents the amino acid sequence of IMX 42 (SEQ ID NO: 35).

Figure 19 presents the amino acid sequence of IMX 44 (SEQ ID NO: 37).

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Figure 20 presents the amino acid sequence of IMX 56 (SEQ ID NO: 38).

Figure 21 presents the nucleotide sequence of the 5' end of the clone (SEQ ID NO:13) matched to part of the human ApoL gene (AF019225).

Figure 22 presents comparison of the 5' end of the clone suggests it represents an alternative splice product to reported ApoL, i.e., bases 1-168 match to 2 exons on PAC carrying the ApoL gene but are not included in the reported complete cDNA.

30

DETAILED DESCRIPTION OF THE INVENTION

Definitions

The following definitions are provided to facilitate understanding of certain terms used
5 throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated"
10 because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a
15 signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence
20 contained in SEQ ID NO:1-26. For example, the polynucleotide can contain all or part of the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the translated amino acid
25 sequence generated from the polynucleotide as broadly defined.

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:1-26, or the complement thereof, or the cDNA. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM
30 NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of

hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH₂PO₄; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

Of course, a polynucleotide which hybridizes only to polyA⁺ sequences (such as any 3' terminal polyA⁺ tract of a cDNA shown in the sequence listing), or to a complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres. and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either

natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formulation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS – STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990); Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.).

The translated amino acid sequence, beginning with the methionine, is identified although other reading frames can also be easily translated using known molecular biology

techniques. The polypeptides produced by the translation of these alternative open reading frames are specifically contemplated by the present invention.

SEQ ID NO:1-26 and the translations of SEQ ID NO: 1-26 as well as SEQ ID NO:27-38 are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from the translations of SEQ ID NO:1-26 may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified.

Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

The present invention also relates to the genes corresponding to SEQ ID NO:1-26, and translations of SEQ ID NO:1-26. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

Also provided in the present invention are species homologues. Species homologues may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for the desired homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below). It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

The polypeptides of the present invention are preferably provided in an isolated form,

and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the invention also can be purified from natural or recombinant sources using antibodies of the invention raised against the secreted protein in methods which are well known in the art.

Signal Sequences

Methods for predicting whether a protein has a signal sequence, as well as the cleavage point for that sequence, are available. For instance, the method of McGeoch, Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 indicates the amino terminus of the secreted protein. Therefore, from a deduced amino acid sequence, a signal sequence and mature sequence can be identified.

Polynucleotide and Polypeptide Variants

"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

Further embodiments of the present invention include polynucleotides having at least 80% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to a sequence contained in SEQ ID NO:1-26. Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the polynucleotides having at least 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity will encode a polypeptide identical to an amino acid sequence contained in the translations of SEQ ID NO:1-26.

Similarly, by a polypeptide having an amino acid sequence having at least, for example, 95% "identity" to a reference polypeptide, is intended that the amino acid sequence of the polypeptide is identical to the reference polypeptide except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the total length of the reference polypeptide. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in

the reference sequence may be deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

Further embodiments of the present invention include polypeptides having at least 80% identity, more preferably at least 85% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to an amino acid sequence contained in translations of SEQ ID NO: 1-26. Preferably, the above polypeptides should exhibit at least one biological activity of the protein.

In a preferred embodiment, polypeptides of the present invention include polypeptides having at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98%, or 99% similarity to an amino acid sequence contained in translations of SEQ ID NO:1-26 as well as the amino acid sequences of SEQ ID NO:27-38.

The variants may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 (1993) reported variant KGF proteins having heparin binding activity even after deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly,

Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)

Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1 α . They used random mutagenesis to generate over 3,500 individual IL-1 α mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." (See Gayle et al., (1993), Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

Thus, the invention further includes polypeptide variants which show substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change. As the authors state, these two strategies, natural selection and genetic engineering, have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile;

replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

5 Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or
10 solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino
15 acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

20

Polynucleotide and Polypeptide Fragments

In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in that shown in SEQ ID NO:1-26. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt,
25 still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in that shown in SEQ ID NO:1-26. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, and more nucleotides) are preferred.

30 Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-260, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, and so forth, to the end of SEQ ID NO:1-26. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably,

these fragments encode a polypeptide which has biological activity.

In the present invention, a "polypeptide fragment" refers to a short amino acid sequence contained in the translations of SEQ ID NO:1-26 as well as SEQ ID NO:27-38. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, and so forth, to the end of the coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50 or 60, amino acids in length. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of the translations of SEQ ID NO:1-26 as well as SEQ ID NO:27-38 falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

Epitopes & Antibodies

In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998-4002 (1983).)

Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')₂ fragments) which are capable of specifically binding to protein. Fab and F(ab')₂ fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred, as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

Fusion Proteins

Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because
5 secreted proteins target cellular locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

10 Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such
15 regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an
20 increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric
25 secreted protein or protein fragment alone. (Fountoulakis et al., J. Biochem. 270:3958-3964 (1995).)

Similarly, EP-A-0 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is
30 beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. (EP-A 0 232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused

with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D. Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. Chem. 270:9459-9471 (1995).)

Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a peptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).)

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

Vectors, Host Cells, and Protein Production

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

Nucleic Acid Molecules and Polypeptides of the Present Invention

The nucleic acid molecules encompassed in the invention comprise nucleotide sequences SEQ ID NO: 1-26.

The amino acid sequences of the polypeptides encoded by the nucleotide sequences of the invention are given in SEQ ID NO:27-38.

The discovery of the nucleic acids of the invention enables the construction of expression vectors comprising nucleic acid sequences encoding polypeptides; host cells transfected or transformed with the expression vectors; isolated and purified biologically active polypeptides and fragments thereof; the use of the nucleic acids or oligonucleotides thereof as probes to identify nucleic acids encoding proteins having amino acid sequences homologous to SEQ ID NOs: 27-38; the use of the nucleic acids or oligonucleotides thereof to identify human chromosomes, for example, 22, 7, and 19; the use of the nucleic acids or oligonucleotides

thereof to map genes on human chromosomes, for example, 22, 7, and 19; the use of the nucleic acids or oligonucleotides thereof to identify genes associated with certain diseases, syndromes or other human conditions associated with human chromosomes such as 22, 7, and 19; the use of single-stranded sense or antisense oligonucleotides from the nucleic acids to inhibit expression of polynucleotides encoded by the IMX sequences; the use of such polypeptides and soluble fragments as molecular weight markers; the use of such polypeptides and fragmented peptides as controls for peptide fragmentation, and kits comprising these reagents; the use of such polypeptides and fragments thereof to generate antibodies, and the use of antibodies to purify IMX polypeptides.

NUCLEIC ACID MOLECULES

In a particular embodiment, the invention relates to certain isolated nucleotide sequences that are free from contaminating endogenous material. A "nucleotide sequence" refers to a polynucleotide molecule in the form of a separate fragment or as a component of a larger nucleic acid construct. The nucleic acid molecule has been derived from DNA or RNA isolated at least once in substantially pure form and in a quantity or concentration enabling identification, manipulation, and recovery of its component nucleotide sequences by standard biochemical methods (such as those outlined in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY (1989)). Such sequences are preferably provided and/or constructed in the form of an open reading frame uninterrupted by internal non-translated sequences, or introns, that are typically present in eukaryotic genes. Sequences of non-translated DNA can be present 5' or 3' from an open reading frame, where the same do not interfere with manipulation or expression of the coding region.

Nucleic acid molecules of the invention include DNA in both single-stranded and double-stranded form, as well as the RNA complement thereof. DNA includes, for example, cDNA, genomic DNA, chemically synthesized DNA, DNA amplified by PCR, and combinations thereof. Genomic DNA may be isolated by conventional techniques, e.g., using the cDNA of SEQ ID NOs:1-26, or a suitable fragment thereof, as a probe.

The DNA molecules of the invention include full length genes as well as polynucleotides and fragments thereof. The full length gene may include the N-terminal signal peptide. Other embodiments include DNA encoding a soluble form, e.g., encoding the extracellular domain of the protein, either with or without the signal peptide.

The nucleic acids of the invention are preferentially derived from human sources, but the invention includes those derived from non-human species, as well.

Preferred Sequences

The particularly preferred nucleotide sequences of the invention are SEQ ID NOs: 1-26, as set forth above. cDNA clones having the nucleotide sequence of SEQ ID NOs:1-26 were
5 isolated as described in Example 1. The sequences of amino acids encoded by the DNA of SEQ ID NOs:1-26 are shown in SEQ ID NOs:27-38.

Additional Sequences

Due to the known degeneracy of the genetic code, wherein more than one codon can
10 encode the same amino acid, a DNA sequence can vary from that shown in SEQ ID NOs:1-26, and still encode a polypeptide having the amino acid sequence of SEQ ID NOs:27-38. Such variant DNA sequences can result from silent mutations (e.g., occurring during PCR amplification), or can be the product of deliberate mutagenesis of a native sequence.

The invention thus provides additional isolated DNA sequences encoding polypeptides
15 of the invention, selected from: (a) DNA comprising the nucleotide sequences of SEQ ID NOs: 1-26; (b) DNA encoding the polypeptides of SEQ ID NOs:27-38; (c) DNA capable of hybridization to a DNA of (a) or (b) under conditions of moderate stringency and which encodes polypeptides of the invention; (d) DNA capable of hybridization to a DNA of (a) or (b) under conditions of high stringency and which encodes polypeptides of the invention, and (e) DNA
20 which is degenerate, as a result of the genetic code, to a DNA defined in (a), (b), (c), or (d) and which encode polypeptides of the invention. Of course, polypeptides encoded by such DNA sequences are encompassed by the invention.

As used herein, conditions of moderate stringency can be readily determined by those having ordinary skill in the art based on, for example, the length of the DNA. The basic
25 conditions are set forth by Sambrook et al. *Molecular Cloning: A Laboratory Manual*, 2 ed. Vol. 1, pp. 1.101-264, Cold Spring Harbor Laboratory Press, (1989), and include use of a prewashing solution for the nitrocellulose filters 5X SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0), hybridization conditions of about 50% formamide, 6X SSC at about 42°C (or other similar hybridization solution, such as Stark's solution, in about 50% formamide at about 42°C), and
30 washing conditions of about 60°C, 0.5X SSC, 0.1% SDS. Conditions of high stringency can also be readily determined by the skilled artisan based on, for example, the length of the DNA. Generally, such conditions are defined as hybridization conditions as above, and with washing at approximately 68°C, 0.2X SSC, 0.1% SDS. The skilled artisan will recognize that the

temperature and wash solution salt concentration can be adjusted as necessary according to factors such as the length of the probe.

Also included as an embodiment of the invention is DNA encoding polypeptide fragments and polypeptides comprising inactivated N-glycosylation site(s), inactivated protease
5 processing site(s), or conservative amino acid substitution(s), as described below.

In another embodiment, the nucleic acid molecules of the invention also comprise nucleotide sequences that are at least 80% identical to a native sequence. Also contemplated are
embodiments in which a nucleic acid molecule comprises a sequence that is at least 90% identical, at least 95% identical, at least 98% identical, at least 99% identical, or at least 99.9%
10 identical to a native sequence.

The percent identity may be determined by visual inspection and mathematical calculation. Alternatively, the percent identity of two nucleic acid sequences can be determined by comparing sequence information using the GAP computer program, version 6.0 described by Devereux et al. (*Nucl. Acids Res.* 12:387, 1984) and available from the University of Wisconsin
15 Genetics Computer Group (UWGCG). The preferred default parameters for the GAP program include: (1) a unary comparison matrix (containing a value of 1 for identities and 0 for non-identities) for nucleotides, and the weighted comparison matrix of Gribskov and Burgess, *Nucl. Acids Res.* 14:6745, 1986, as described by Schwartz and Dayhoff, eds., *Atlas of Protein Sequence and Structure*, National Biomedical Research Foundation, pp. 353-358, 1979; (2) a
20 penalty of 3.0 for each gap and an additional 0.10 penalty for each symbol in each gap; and (3) no penalty for end gaps. Other programs used by one skilled in the art of sequence comparison may also be used.

The invention provides isolated nucleic acids useful in the production of polypeptides. Such polypeptides may be prepared by any of a number of conventional techniques. A DNA
25 sequence encoding an IMX polypeptide or a desired fragment thereof may be subcloned into an expression vector for production of the polypeptide or fragment. The DNA sequence advantageously is fused to a sequence encoding a suitable leader or signal peptide. Alternatively, the desired fragment may be chemically synthesized using known techniques. DNA fragments also may be produced by restriction endonuclease digestion of a full length
30 cloned DNA sequence, and isolated by electrophoresis on agarose gels. If necessary, oligonucleotides that reconstruct the 5' or 3' terminus to a desired point may be ligated to a DNA fragment generated by restriction enzyme digestion. Such oligonucleotides may additionally contain a restriction endonuclease cleavage site upstream of the desired coding sequence, and position an initiation codon (ATG) at the N-terminus of the coding sequence.

The well-known polymerase chain reaction (PCR) procedure also may be employed to isolate and amplify a DNA sequence encoding a desired protein fragment. Oligonucleotides that define the desired termini of the DNA fragment are employed as 5' and 3' primers. The oligonucleotides may additionally contain recognition sites for restriction endonucleases, to facilitate insertion of the amplified DNA fragment into an expression vector. PCR techniques are described in Saiki et al., *Science* 239:487 (1988); *Recombinant DNA Methodology*, Wu et al., eds., Academic Press, Inc., San Diego (1989), pp. 189-196; and *PCR Protocols: A Guide to Methods and Applications*, Innis et al., eds., Academic Press, Inc. (1990).

10 POLYPEPTIDES AND FRAGMENTS THEREOF

The invention encompasses polypeptides and fragments thereof in various forms, including those that are naturally occurring or produced through various techniques such as procedures involving recombinant DNA technology. Such forms include, but are not limited to, derivatives, variants, and oligomers, as well as fusion proteins or fragments thereof.

15 Polypeptides and Fragments Thereof

The polypeptides of the invention include full length proteins encoded by the nucleic acid sequences set forth above. Particularly preferred polypeptides comprise the amino acid sequences of SEQ ID NOs:27-38.

20 The polypeptides of the invention may include an N-terminal hydrophobic region that functions as a signal peptide, and may contain an extracellular domain, and may also contain a transmembrane region and a C-terminal cytoplasmic domain as well as a spacer region. Computer analysis may be used to predict the location of the signal peptide.

For example, the isolated polypeptides of SEQ ID NO:14 (IMX 28) and SEQ ID NO:19 (IMX 44) include an N-terminal hydrophobic region that functions as a signal peptide. Computer analysis predicts that the signal peptide corresponds to residues 1 to 17 of SEQ ID NO:14, and cleavage of the signal peptide of SEQ ID NO:14 results in a mature protein comprising amino acids 18 to 372. The signal peptide of IMX 44 corresponds to residues 1 to 37 of SEQ ID NO:19. The next most likely computer-predicted signal peptide cleavage sites (in descending order) would occur after amino acids 36, 26 and 27 of SEQ ID NO:19. Cleavage of the signal peptide at position 37 thus would yield a mature protein comprising amino acids 38 through 261 of SEQ ID NO:19.

The skilled artisan will recognize that the above-described boundaries of such regions of the polypeptide are approximate. To illustrate, the boundaries of the mature protein (which may

be predicted by using computer programs available for that purpose) may differ from those described above.

The polypeptides of the invention may be membrane bound or they may be secreted and, thus, soluble. Soluble polypeptides are capable of being secreted from the cells in which they are expressed. In general, soluble polypeptides may be identified (and distinguished from non-soluble membrane-bound counterparts) by separating intact cells which express the desired polypeptide from the culture medium, *e.g.*, by centrifugation, and assaying the medium (supernatant) for the presence of the desired polypeptide. The presence of polypeptide in the medium indicates that the polypeptide was secreted from the cells and thus is a soluble form of the protein.

In one embodiment, the soluble polypeptides and fragments thereof comprise all or part of the extracellular domain, but lack the transmembrane region that would cause retention of the polypeptide on a cell membrane. A soluble polypeptide may include the cytoplasmic domain, or a portion thereof, as long as the polypeptide is secreted from the cell in which it is produced.

In general, the use of soluble forms is advantageous for certain applications. Purification of the polypeptides from recombinant host cells is facilitated, since the soluble polypeptides are secreted from the cells. Further, soluble polypeptides are generally more suitable for intravenous administration.

The invention also provides polypeptides and fragments of the extracellular domain that retain a desired biological activity. Particular embodiments are directed to polypeptide fragments that retain the ability to bind the native cognate, substrate, or counter-structure ("binding partner"). Such a fragment may be a soluble polypeptide, as described above. In another embodiment, the polypeptides and fragments advantageously include regions that are conserved in the family as described above.

Also provided herein are polypeptide fragments comprising at least 20, or at least 30, contiguous amino acids of the sequence of SEQ ID NOs:27-38. Fragments derived from the cytoplasmic domain find use in studies of signal transduction, and in regulating cellular processes associated with transduction of biological signals. Polypeptide fragments also may be employed as immunogens, in generating antibodies.

Variants

Naturally occurring variants as well as derived variants of the polypeptides and fragments are provided herein.

Variants may exhibit amino acid sequences that are at least 80% identical. Also contemplated are embodiments in which a polypeptide or fragment comprises an amino acid sequence that is at least 90% identical, at least 95% identical, at least 98% identical, at least 99% identical, or at least 99.9% identical to the preferred polypeptide or fragment thereof. Percent identity may be determined by visual inspection and mathematical calculation. Alternatively, the percent identity of two protein sequences can be determined by comparing sequence information using the GAP computer program, based on the algorithm of Needleman and Wunsch (J. Mol. Bio. 48:443, 1970) and available from the University of Wisconsin Genetics Computer Group (UWGCG). The preferred default parameters for the GAP program include: (1) a scoring matrix, blosum62, as described by Henikoff and Henikoff (Proc. Natl. Acad. Sci. USA 89:10915, 1992); (2) a gap weight of 12; (3) a gap length weight of 4; and (4) no penalty for end gaps. Other programs used by one skilled in the art of sequence comparison may also be used.

The variants of the invention include, for example, those that result from alternate mRNA splicing events or from proteolytic cleavage. Alternate splicing of mRNA may, for example, yield a truncated but biologically active protein, such as a naturally occurring soluble form of the protein. Variations attributable to proteolysis include, for example, differences in the N- or C-termini upon expression in different types of host cells, due to proteolytic removal of one or more terminal amino acids from the protein (generally from 1-5 terminal amino acids). Proteins in which differences in amino acid sequence are attributable to genetic polymorphism (allelic variation among individuals producing the protein) are also contemplated herein.

Additional variants within the scope of the invention include polypeptides that may be modified to create derivatives thereof by forming covalent or aggregative conjugates with other chemical moieties, such as glycosyl groups, lipids, phosphate, acetyl groups and the like. Covalent derivatives may be prepared by linking the chemical moieties to functional groups on amino acid side chains or at the N-terminus or C-terminus of a polypeptide. Conjugates comprising diagnostic (detectable) or therapeutic agents attached thereto are contemplated herein, as discussed in more detail below.

Other derivatives include covalent or aggregative conjugates of the polypeptides with other proteins or polypeptides, such as by synthesis in recombinant culture as N-terminal or C-terminal fusions. Examples of fusion proteins are discussed below in connection with oligomers. Further, fusion proteins can comprise peptides added to facilitate purification and identification. Such peptides include, for example, poly-His or the antigenic identification peptides described in U.S. Patent No. 5,011,912 and in Hopp et al., *Bio/Technology* 6:1204,

1988. One such peptide is the FLAG[®] peptide, Asp-Tyr-Lys-Asp-Asp-Asp-Lys, which is highly antigenic and provides an epitope reversibly bound by a specific monoclonal antibody, enabling rapid assay and facile purification of expressed recombinant protein. A murine hybridoma designated 4E11 produces a monoclonal antibody that binds the FLAG[®] peptide in the presence of certain divalent metal cations, as described in U.S. Patent 5,011,912, hereby incorporated by reference. The 4E11 hybridoma cell line has been deposited with the American Type Culture Collection under accession no. HB 9259. Monoclonal antibodies that bind the FLAG[®] peptide are available from Eastman Kodak Co., Scientific Imaging Systems Division, New Haven, Connecticut.

Among the polypeptides provided herein are variants of native polypeptides that retain the native biological activity or the substantial equivalent thereof. One example is a variant that binds with essentially the same binding affinity as does the native form. Binding affinity can be measured by conventional procedures, *e.g.*, as described in U.S. Patent No. 5,512,457 and as set forth below.

Variants include polypeptides that are substantially homologous to the native form, but which have an amino acid sequence different from that of the native form because of one or more deletions, insertions or substitutions. Particular embodiments include, but are not limited to, polypeptides that comprise from one to ten deletions, insertions or substitutions of amino acid residues, when compared to a native sequence.

A given amino acid may be replaced, for example, by a residue having similar physiochemical characteristics. Examples of such conservative substitutions include substitution of one aliphatic residue for another, such as Ile, Val, Leu, or Ala for one another; substitutions of one polar residue for another, such as between Lys and Arg, Glu and Asp, or Gln and Asn; or substitutions of one aromatic residue for another, such as Phe, Trp, or Tyr for one another. Other conservative substitutions, *e.g.*, involving substitutions of entire regions having similar hydrophobicity characteristics, are well known.

Similarly, the DNAs of the invention include variants that differ from a native DNA sequence because of one or more deletions, insertions or substitutions, but that encode a biologically active polypeptide.

The invention further includes polypeptides of the invention with or without associated native-pattern glycosylation. Polypeptides expressed in yeast or mammalian expression systems (*e.g.*, COS-1 or COS-7 cells) can be similar to or significantly different from a native polypeptide in molecular weight and glycosylation pattern, depending upon the choice of expression system. Expression of polypeptides of the invention in bacterial expression systems.

such as *E. coli*, provides non-glycosylated molecules. Further, a given preparation may include multiple differentially glycosylated species of the protein. Glycosyl groups can be removed through conventional methods, in particular those utilizing glycopeptidase. In general, glycosylated polypeptides of the invention can be incubated with a molar excess of glycopeptidase (Boehringer Mannheim).

Correspondingly, similar DNA constructs that encode various additions or substitutions of amino acid residues or sequences, or deletions of terminal or internal residues or sequences are encompassed by the invention. For example, N-glycosylation sites in the polypeptide extracellular domain can be modified to preclude glycosylation, allowing expression of a reduced carbohydrate analog in mammalian and yeast expression systems. N-glycosylation sites in eukaryotic polypeptides are characterized by an amino acid triplet Asn-X-Y, wherein X is any amino acid except Pro and Y is Ser or Thr. Appropriate substitutions, additions, or deletions to the nucleotide sequence encoding these triplets will result in prevention of attachment of carbohydrate residues at the Asn side chain. Alteration of a single nucleotide, chosen so that Asn is replaced by a different amino acid, for example, is sufficient to inactivate an N-glycosylation site. Alternatively, the Ser or Thr can be replaced with another amino acid, such as Ala. Known procedures for inactivating N-glycosylation sites in proteins include those described in U.S. Patent 5,071,972 and EP 276,846, hereby incorporated by reference.

In another example of variants, sequences encoding Cys residues that are not essential for biological activity can be altered to cause the Cys residues to be deleted or replaced with other amino acids, preventing formation of incorrect intramolecular disulfide bridges upon folding or renaturation.

Other variants are prepared by modification of adjacent dibasic amino acid residues, to enhance expression in yeast systems in which KEX2 protease activity is present. EP 212,914 discloses the use of site-specific mutagenesis to inactivate KEX2 protease processing sites in a protein. KEX2 protease processing sites are inactivated by deleting, adding or substituting residues to alter Arg-Arg, Arg-Lys, and Lys-Arg pairs to eliminate the occurrence of these adjacent basic residues. Lys-Lys pairings are considerably less susceptible to KEX2 cleavage, and conversion of Arg-Lys or Lys-Arg to Lys-Lys represents a conservative and preferred approach to inactivating KEX2 sites.

Oligomers

Encompassed by the invention are oligomers or fusion proteins that contain IMX polypeptides. Such oligomers may be in the form of covalently-linked or non-covalently-linked multimers, including dimers, trimers, or higher oligomers. As noted above, preferred
5 polypeptides are soluble and thus these oligomers may comprise soluble polypeptides. In one aspect of the invention, the oligomers maintain the binding ability of the polypeptide components and provide therefor, bivalent, trivalent, etc., binding sites.

One embodiment of the invention is directed to oligomers comprising multiple polypeptides joined *via* covalent or non-covalent interactions between peptide moieties fused to
10 the polypeptides. Such peptides may be peptide linkers (spacers), or peptides that have the property of promoting oligomerization. Leucine zippers and certain polypeptides derived from antibodies are among the peptides that can promote oligomerization of the polypeptides attached thereto, as described in more detail below.

Immunoglobulin-based Oligomers

15 As one alternative, an oligomer is prepared using polypeptides derived from immunoglobulins. Preparation of fusion proteins comprising certain heterologous polypeptides fused to various portions of antibody-derived polypeptides (including the Fc domain) has been described, *e.g.*, by Ashkenazi et al. (*PNAS USA* 88:10535, 1991); Byrn et al. (*Nature* 344:677, 1990); and Hollenbaugh and Aruffo ("Construction of Immunoglobulin Fusion Proteins", in
20 *Current Protocols in Immunology*, Suppl. 4, pages 10.19.1 - 10.19.11, 1992).

One embodiment of the present invention is directed to a dimer comprising two fusion proteins created by fusing a polypeptide of the invention to an Fc polypeptide derived from an antibody. A gene fusion encoding the polypeptide/Fc fusion protein is inserted into an appropriate expression vector. Polypeptide/Fc fusion proteins are expressed in host cells
25 transformed with the recombinant expression vector, and allowed to assemble much like antibody molecules, whereupon interchain disulfide bonds form between the Fc moieties to yield divalent molecules.

The term "Fc polypeptide" as used herein includes native and mutein forms of polypeptides comprising from the Fc region of an antibody. Truncated forms of such
30 polypeptides containing the hinge region that promotes dimerization are also included. Preferred polypeptides comprise an Fc polypeptide derived from a human IgG1 antibody comprising any or all of the CH domains of the Fc region.

One suitable Fc polypeptide, described in PCT application WO 93/10151, hereby incorporated by reference, is a single chain polypeptide extending from the N-terminal hinge

region to the native C-terminus of the Fc region of a human IgG1 antibody. Another useful Fc polypeptide is the Fc mutein described in U.S. Patent 5,457,035 and in Baum et al., (*EMBO J.* 13:3992-4001, 1994) incorporated herein by reference. The amino acid sequence of this mutein is identical to that of the native Fc sequence presented in WO 93/10151, except that amino acid
5 19 has been changed from Leu to Ala, amino acid 20 has been changed from Leu to Glu, and amino acid 22 has been changed from Gly to Ala. The mutein exhibits reduced affinity for Fc receptors.

The above-described fusion proteins comprising Fc moieties (and oligomers formed therefrom) offer the advantage of facile purification by affinity chromatography over Protein A
10 or Protein G columns.

In other embodiments, the polypeptides of the invention may be substituted for the variable portion of an antibody heavy or light chain. If fusion proteins are made with both heavy and light chains of an antibody, it is possible to form an oligomer with as many as four soluble regions of the proteins of the invention.

Peptide-linker Based Oligomers

Alternatively, the oligomer is a fusion protein comprising multiple polypeptides, with or without peptide linkers (spacer peptides). Among the suitable peptide linkers are those described in U.S. Patents 4,751,180 and 4,935,233, which are hereby incorporated by reference.
20 A DNA sequence encoding a desired peptide linker may be inserted between, and in the same reading frame as, the DNA sequences of the invention, using any suitable conventional technique. For example, a chemically synthesized oligonucleotide encoding the linker may be ligated between the sequences. In particular embodiments, a fusion protein comprises from two to four soluble IMX polypeptides, separated by peptide linkers.

Leucine-Zippers

Another method for preparing the oligomers of the invention involves use of a leucine zipper. Leucine zipper domains are peptides that promote oligomerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding
30 proteins (Landschulz et al., *Science* 240:1759, 1988), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize.

Examples of leucine zipper domains suitable for producing soluble oligomeric proteins are described in PCT application WO 94/10308, and the leucine zipper derived from lung

surfactant protein D (SPD) described in Hoppe et al. (*FEBS Letters* 344:191, 1994), hereby incorporated by reference. The use of a modified leucine zipper that allows for stable trimerization of a heterologous protein fused thereto is described in Fanslow et al. (*Semin. Immunol.* 6:267-278, 1994). Recombinant fusion proteins comprising a soluble polypeptide fused to a leucine zipper peptide are expressed in suitable host cells, and the soluble oligomer that forms is recovered from the culture supernatant.

In particular embodiments, leucine residues in a leucine zipper moiety are replaced by isoleucine residues. Such peptides comprising isoleucine may be referred to as "isoleucine zippers" but are encompassed by the term "leucine zippers" as employed herein.

PRODUCTION OF POLYPEPTIDES AND FRAGMENTS THEREOF

Expression, isolation and purification of the polypeptides and fragments of the invention may be accomplished by any suitable technique, including but not limited to the following:

Expression Systems

The present invention also provides recombinant cloning and expression vectors containing DNA, as well as host cell containing the recombinant vectors. Expression vectors comprising DNA may be used to prepare the polypeptides or fragments of the invention encoded by the DNA. A method for producing polypeptides comprises culturing host cells transformed with a recombinant expression vector encoding the polypeptide, under conditions that promote expression of the polypeptide, then recovering the expressed polypeptides from the culture. The skilled artisan will recognize that the procedure for purifying the expressed polypeptides will vary according to such factors as the type of host cells employed, and whether the polypeptide is membrane-bound or a soluble form that is secreted from the host cell.

Any suitable expression system may be employed. The vectors include a DNA encoding a polypeptide or fragment of the invention, operably linked to suitable transcriptional or translational regulatory nucleotide sequences, such as those derived from a mammalian, microbial, viral, or insect gene. Examples of regulatory sequences include transcriptional promoters, operators, or enhancers, an mRNA ribosomal binding site, and appropriate sequences which control transcription and translation initiation and termination. Nucleotide sequences are operably linked when the regulatory sequence functionally relates to the DNA sequence. Thus, a promoter nucleotide sequence is operably linked to a DNA sequence if the promoter nucleotide sequence controls the transcription of the DNA sequence. An origin of replication that confers

the ability to replicate in the desired host cells, and a selection gene by which transformants are identified, are generally incorporated into the expression vector.

In addition, a sequence encoding an appropriate signal peptide (native or heterologous) can be incorporated into expression vectors. A DNA sequence for a signal peptide (secretory leader) may be fused in frame to the nucleic acid sequence of the invention so that the DNA is initially transcribed, and the mRNA translated, into a fusion protein comprising the signal peptide. A signal peptide that is functional in the intended host cells promotes extracellular secretion of the polypeptide. The signal peptide is cleaved from the polypeptide upon secretion of polypeptide from the cell.

The skilled artisan will also recognize that the position(s) at which the signal peptide is cleaved may differ from that predicted by computer program, and may vary according to such factors as the type of host cells employed in expressing a recombinant polypeptide. A protein preparation may include a mixture of protein molecules having different N-terminal amino acids, resulting from cleavage of the signal peptide at more than one site.

Suitable host cells for expression of polypeptides include prokaryotes, yeast or higher eukaryotic cells. Mammalian or insect cells are generally preferred for use as host cells. Appropriate cloning and expression vectors for use with bacterial, fungal, yeast, and mammalian cellular hosts are described, for example, in Pouwels et al. *Cloning Vectors: A Laboratory Manual*, Elsevier, New York, (1985). Cell-free translation systems could also be employed to produce polypeptides using RNAs derived from DNA constructs disclosed herein.

Prokaryotic Systems

Prokaryotes include gram-negative or gram-positive organisms. Suitable prokaryotic host cells for transformation include, for example, *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium*, and various other species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*. In a prokaryotic host cell, such as *E. coli*, a polypeptide may include an N-terminal methionine residue to facilitate expression of the recombinant polypeptide in the prokaryotic host cell. The N-terminal Met may be cleaved from the expressed recombinant polypeptide.

Expression vectors for use in prokaryotic host cells generally comprise one or more phenotypic selectable marker genes. A phenotypic selectable marker gene is, for example, a gene encoding a protein that confers antibiotic resistance or that supplies an autotrophic requirement. Examples of useful expression vectors for prokaryotic host cells include those derived from commercially available plasmids such as the cloning vector pBR322 (ATCC

37017). pBR322 contains genes for ampicillin and tetracycline resistance and thus provides simple means for identifying transformed cells. An appropriate promoter and a DNA sequence are inserted into the pBR322 vector. Other commercially available vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and pGEM1 (Promega Biotec,
5 Madison, WI, USA).

Promoter sequences commonly used for recombinant prokaryotic host cell expression vectors include β -lactamase (penicillinase), lactose promoter system (Chang et al., *Nature* 275:615, 1978; and Goeddel et al., *Nature* 281:544, 1979), tryptophan (trp) promoter system (Goeddel et al., *Nucl. Acids Res.* 8:4057, 1980; and EP-A-36776) and tac promoter (Maniatis,
10 *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory, p. 412, 1982). A particularly useful prokaryotic host cell expression system employs a phage λP_L promoter and a cI857ts thermolabile repressor sequence. Plasmid vectors available from the American Type Culture Collection which incorporate derivatives of the λP_L promoter include plasmid pHUB2 (resident in *E. coli* strain JMB9, ATCC 37092) and pPLc28 (resident in *E. coli* RR1, ATCC
15 53082).

Yeast Systems

Alternatively, the polypeptides may be expressed in yeast host cells, preferably from the *Saccharomyces* genus (e.g., *S. cerevisiae*). Other genera of yeast, such as *Pichia* or
20 *Kluyveromyces*, may also be employed. Yeast vectors will often contain an origin of replication sequence from a 2 μ yeast plasmid, an autonomously replicating sequence (ARS), a promoter region, sequences for polyadenylation, sequences for transcription termination, and a selectable marker gene. Suitable promoter sequences for yeast vectors include, among others, promoters for metallothionein, 3-phosphoglycerate kinase (Hitzeman et al., *J. Biol. Chem.* 255:2073, 1980)
25 or other glycolytic enzymes (Hess et al., *J. Adv. Enzyme Reg.* 7:149, 1968; and Holland et al., *Biochem.* 17:4900, 1978), such as enolase, glyceraldehyde-3-phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phosphofructokinase, glucose-6-phosphate isomerase, 3-phosphoglycerate mutase, pyruvate kinase, triosephosphate isomerase, phospho-glucose isomerase, and glucokinase. Other suitable vectors and promoters for use in yeast expression
30 are further described in Hitzeman, EPA-73,657. Another alternative is the glucose-repressible ADH2 promoter described by Russell et al. (*J. Biol. Chem.* 258:2674, 1982) and Beier et al. (*Nature* 300:724, 1982). Shuttle vectors replicable in both yeast and *E. coli* may be constructed by inserting DNA sequences from pBR322 for selection and replication in *E. coli* (Amp^r gene and origin of replication) into the above-described yeast vectors.

The yeast α -factor leader sequence may be employed to direct secretion of the polypeptide. The α -factor leader sequence is often inserted between the promoter sequence and the structural gene sequence. See, e.g., Kurjan et al., *Cell* 30:933, 1982 and Bitter et al., *Proc. Natl. Acad. Sci. USA* 81:5330, 1984. Other leader sequences suitable for facilitating secretion of recombinant polypeptides from yeast hosts are known to those of skill in the art. A leader sequence may be modified near its 3' end to contain one or more restriction sites. This will facilitate fusion of the leader sequence to the structural gene.

Yeast transformation protocols are known to those of skill in the art. One such protocol is described by Hinnen et al., *Proc. Natl. Acad. Sci. USA* 75:1929, 1978. The Hinnen et al. protocol selects for Trp⁺ transformants in a selective medium, wherein the selective medium consists of 0.67% yeast nitrogen base, 0.5% casamino acids, 2% glucose, 10 mg/ml adenine and 20 mg/ml uracil.

Yeast host cells transformed by vectors containing an ADH2 promoter sequence may be grown for inducing expression in a "rich" medium. An example of a rich medium is one consisting of 1% yeast extract, 2% peptone, and 1% glucose supplemented with 80 mg/ml adenine and 80 mg/ml uracil. Derepression of the ADH2 promoter occurs when glucose is exhausted from the medium.

Mammalian or Insect Systems

Mammalian or insect host cell culture systems also may be employed to express recombinant polypeptides. Bacculovirus systems for production of heterologous proteins in insect cells are reviewed by Luckow and Summers, *Bio/Technology* 6:47 (1988). Established cell lines of mammalian origin also may be employed. Examples of suitable mammalian host cell lines include the COS-7 line of monkey kidney cells (ATCC CRL 1651) (Gluzman et al., *Cell* 23:175, 1981), L cells, C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells, HeLa cells, and BHK (ATCC CRL 10) cell lines, and the CV1/EBNA cell line derived from the African green monkey kidney cell line CV1 (ATCC CCL 70) as described by McMahan et al. (*EMBO J.* 10: 2821, 1991).

Established methods for introducing DNA into mammalian cells have been described (Kaufman, R.J., *Large Scale Mammalian Cell Culture*, 1990, pp. 15-69). Additional protocols using commercially available reagents, such as Lipofectamine lipid reagent (Gibco/BRL) or Lipofectamine-Plus lipid reagent, can be used to transfect cells (Felgner et al., *Proc. Natl. Acad. Sci. USA* 84:7413-7417, 1987). In addition, electroporation can be used to transfect mammalian cells using conventional procedures, such as those in Sambrook et al. (*Molecular Cloning: A*

Laboratory Manual, 2 ed. Vol. 1-3, Cold Spring Harbor Laboratory Press, 1989). Selection of stable transformants can be performed using methods known in the art, such as, for example, resistance to cytotoxic drugs. Kaufman et al., *Meth. in Enzymology* 185:487-511, 1990, describes several selection schemes, such as dihydrofolate reductase (DHFR) resistance. A suitable host strain for DHFR selection can be CHO strain DX-B11, which is deficient in DHFR (Urlaub and Chasin, *Proc. Natl. Acad. Sci. USA* 77:4216-4220, 1980). A plasmid expressing the DHFR cDNA can be introduced into strain DX-B11, and only cells that contain the plasmid can grow in the appropriate selective media. Other examples of selectable markers that can be incorporated into an expression vector include cDNAs conferring resistance to antibiotics, such as G418 and hygromycin B. Cells harboring the vector can be selected on the basis of resistance to these compounds.

Transcriptional and translational control sequences for mammalian host cell expression vectors can be excised from viral genomes. Commonly used promoter sequences and enhancer sequences are derived from polyoma virus, adenovirus 2, simian virus 40 (SV40), and human cytomegalovirus. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early and late promoter, enhancer, splice, and polyadenylation sites can be used to provide other genetic elements for expression of a structural gene sequence in a mammalian host cell. Viral early and late promoters are particularly useful because both are easily obtained from a viral genome as a fragment, which can also contain a viral origin of replication (Fiers et al., *Nature* 273:113, 1978; Kaufman, *Meth. in Enzymology*, 1990). Smaller or larger SV40 fragments can also be used, provided the approximately 250 bp sequence extending from the *Hind* III site toward the *Bgl* I site located in the SV40 viral origin of replication site is included.

Additional control sequences shown to improve expression of heterologous genes from mammalian expression vectors include such elements as the expression augmenting sequence element (EASE) derived from CHO cells (Morris et al., *Animal Cell Technology*, 1997, pp. 529-534 and PCT Application WO 97/25420) and the tripartite leader (TPL) and VA gene RNAs from Adenovirus 2 (Gingeras et al., *J. Biol. Chem.* 257:13475-13491, 1982). The internal ribosome entry site (IRES) sequences of viral origin allows dicistronic mRNAs to be translated efficiently (Oh and Sarnow, *Current Opinion in Genetics and Development* 3:295-300, 1993; Ramesh et al., *Nucleic Acids Research* 24:2697-2700, 1996). Expression of a heterologous cDNA as part of a dicistronic mRNA followed by the gene for a selectable marker (e.g. DHFR) has been shown to improve transfectability of the host and expression of the heterologous cDNA (Kaufman, *Meth. in Enzymology*, 1990). Exemplary expression vectors that employ dicistronic

mRNAs are pTR-DC/GFP described by Mosser et al., *Biotechniques* 22:150-161, 1997, and p2A5I described by Morris et al., *Animal Cell Technology*, 1997, pp. 529-534.

5 A useful high expression vector, pCAVNOT, has been described by Mosley et al., *Cell* 59:335-348, 1989. Other expression vectors for use in mammalian host cells can be constructed as disclosed by Okayama and Berg (*Mol. Cell. Biol.* 3:280, 1983). A useful system for stable high level expression of mammalian cDNAs in C127 murine mammary epithelial cells can be constructed substantially as described by Cosman et al. (*Mol. Immunol.* 23:935, 1986). A useful high expression vector, PMLSV N1/N4, described by Cosman et al., *Nature* 312:768, 1984, has been deposited as ATCC 39890. Additional useful mammalian expression vectors are described
10 in EP-A-0367566, and in WO 91/18982, incorporated by reference herein. In yet another alternative, the vectors can be derived from retroviruses.

Another useful expression vector, pFLAG[®], can be used. FLAG[®] technology is centered on the fusion of a low molecular weight (1kD), hydrophilic, FLAG[®] marker peptide to the N-terminus of a recombinant protein expressed by pFLAG[®] expression vectors.

15 Regarding signal peptides that may be employed, the native signal peptide may be replaced by a heterologous signal peptide or leader sequence, if desired. The choice of signal peptide or leader may depend on factors such as the type of host cells in which the recombinant polypeptide is to be produced. To illustrate, examples of heterologous signal peptides that are functional in mammalian host cells include the signal sequence for interleukin-7 (IL-7)
20 described in United States Patent 4,965,195, the signal sequence for interleukin-2 receptor described in Cosman et al., *Nature* 312:768 (1984); the interleukin-4 receptor signal peptide described in EP 367,566; the type I interleukin-1 receptor signal peptide described in U.S. Patent 4,968,607; and the type II interleukin-1 receptor signal peptide described in EP 460,846.

Purification

25 The invention also includes methods of isolating and purifying the polypeptides and fragments thereof.

Isolation and Purification

The "isolated" polypeptides or fragments thereof encompassed by this invention are polypeptides or fragments that are not in an environment identical to an environment in which it or they can be found in nature. The "purified" polypeptides or fragments thereof encompassed
5 by this invention are essentially free of association with other proteins or polypeptides, for example, as a purification product of recombinant expression systems such as those described above or as a purified product from a non-recombinant source such as naturally occurring cells and/or tissues.

In one preferred embodiment, the purification of recombinant polypeptides or fragments
10 can be accomplished using fusions of polypeptides or fragments of the invention to another polypeptide to aid in the purification of polypeptides or fragments of the invention. Such fusion partners can include the poly-His or other antigenic identification peptides described above as well as the Fc moieties described previously.

With respect to any type of host cell, as is known to the skilled artisan, procedures for
15 purifying a recombinant polypeptide or fragment will vary according to such factors as the type of host cells employed and whether or not the recombinant polypeptide or fragment is secreted into the culture medium.

In general, the recombinant polypeptide or fragment can be isolated from the host cells if not secreted, or from the medium or supernatant if soluble and secreted, followed by one or
20 more concentration, salting-out, ion exchange, hydrophobic interaction, affinity purification or size exclusion chromatography steps. As to specific ways to accomplish these steps, the culture medium first can be concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. Following the concentration step, the concentrate can be applied to a purification matrix such as a gel filtration medium.
25 Alternatively, an anion exchange resin can be employed, for example, a matrix or substrate having pendant diethylaminoethyl (DEAE) groups. The matrices can be acrylamide, agarose, dextran, cellulose or other types commonly employed in protein purification. Alternatively, a cation exchange step can be employed. Suitable cation exchangers include various insoluble matrices comprising sulfopropyl or carboxymethyl groups. In addition, a chromatofocusing step
30 can be employed. Alternatively, a hydrophobic interaction chromatography step can be employed. Suitable matrices can be phenyl or octyl moieties bound to resins. In addition, affinity chromatography with a matrix which selectively binds the recombinant protein can be employed. Examples of such resins employed are lectin columns, dye columns, and metal-chelating columns. Finally, one or more reverse-phase high performance liquid chromatography

(RP-HPLC) steps employing hydrophobic RP-HPLC media. (e.g., silica gel or polymer resin having pendant methyl, octyl, octyldecyl or other aliphatic groups) can be employed to further purify the polypeptides. Some or all of the foregoing purification steps, in various combinations, are well known and can be employed to provide an isolated and purified recombinant protein.

It is also possible to utilize an affinity column comprising a polypeptide-binding protein of the invention, such as a monoclonal antibody generated against polypeptides of the invention, to affinity-purify expressed polypeptides. These polypeptides can be removed from an affinity column using conventional techniques, e.g., in a high salt elution buffer and then dialyzed into a lower salt buffer for use or by changing pH or other components depending on the affinity matrix utilized, or be competitively removed using the naturally occurring substrate of the affinity moiety, such as a polypeptide derived from the invention.

In this aspect of the invention, polypeptide-binding proteins, such as the anti-polypeptide antibodies of the invention or other proteins that may interact with the polypeptide of the invention, can be bound to a solid phase support such as a column chromatography matrix or a similar substrate suitable for identifying, separating, or purifying cells that express polypeptides of the invention on their surface. Adherence of polypeptide-binding proteins of the invention to a solid phase contacting surface can be accomplished by any means. For example, magnetic microspheres can be coated with these polypeptide-binding proteins and held in the incubation vessel through a magnetic field. Suspensions of cell mixtures are contacted with the solid phase that has such polypeptide-binding proteins thereon. Cells having polypeptides of the invention on their surface bind to the fixed polypeptide-binding protein and unbound cells then are washed away. This affinity-binding method is useful for purifying, screening, or separating such polypeptide-expressing cells from solution. Methods of releasing positively selected cells from the solid phase are known in the art and encompass, for example, the use of enzymes. Such enzymes are preferably non-toxic and non-injurious to the cells and are preferably directed to cleaving the cell-surface binding partner.

Alternatively, mixtures of cells suspected of containing polypeptide-expressing cells of the invention first can be incubated with a biotinylated polypeptide-binding protein of the invention. Incubation periods are typically at least one hour in duration to ensure sufficient binding to polypeptides of the invention. The resulting mixture then is passed through a column packed with avidin-coated beads, whereby the high affinity of biotin for avidin provides the binding of the polypeptide-binding cells to the beads. Use of avidin-coated beads is known in

the art. See Berenson, et al. *J. Cell. Biochem.*, 10D:239 (1986). Wash of unbound material and the release of the bound cells is performed using conventional methods.

The desired degree of purity depends on the intended use of the protein. A relatively high degree of purity is desired when the polypeptide is to be administered *in vivo*, for example. In such a case, the polypeptides are purified such that no protein bands corresponding to other proteins are detectable upon analysis by SDS-polyacrylamide gel electrophoresis (SDS-PAGE). It will be recognized by one skilled in the pertinent field that multiple bands corresponding to the polypeptide may be visualized by SDS-PAGE, due to differential glycosylation, differential post-translational processing, and the like. Most preferably, the polypeptide of the invention is purified to substantial homogeneity, as indicated by a single protein band upon analysis by SDS-PAGE. The protein band may be visualized by silver staining, Coomassie blue staining, or (if the protein is radiolabeled) by autoradiography.

Assays

The purified polypeptides of the invention (including proteins, polypeptides, fragments, variants, oligomers, and other forms) may be tested for the ability to bind a cognate, ligand, receptor, substrate, or counter-structure and the like ("binding partner") in any suitable assay, such as a conventional binding assay. To illustrate, the polypeptide may be labeled with a detectable reagent (e.g., a radionuclide, chromophore, enzyme that catalyzes a colorimetric or fluorometric reaction, and the like). The labeled polypeptide is contacted with cells expressing the binding partner. The cells then are washed to remove unbound labeled polypeptide, and the presence of cell-bound label is determined by a suitable technique, chosen according to the nature of the label.

One example of a binding assay procedure is as follows. A recombinant expression vector containing the binding partner cDNA is constructed using methods known in the art. CV1-EBNA-1 cells in 10 cm² dishes are transfected with the recombinant expression vector. CV-1/EBNA-1 cells (ATCC CRL 10478) constitutively express EBV nuclear antigen-1 driven from the CMV immediate-early enhancer/promoter. CV1-EBNA-1 was derived from the African Green Monkey kidney cell line CV-1 (ATCC CCL 70), as described by McMahan et al. (*EMBO J.* 10:2821, 1991).

The transfected cells are cultured for 24 hours, and the cells in each dish then are split into a 24-well plate. After culturing an additional 48 hours, the transfected cells (about 4 x 10⁴ cells/well) are washed with BM-NFDM, which is binding medium (RPMI 1640 containing 25 mg/ml bovine serum albumin, 2 mg/ml sodium azide, 20 mM Hepes pH 7.2) to which 50 mg/ml

nonfat dry milk has been added. The cells then are incubated for 1 hour at 37°C with various concentrations of, for example, a soluble polypeptide/Fc fusion protein made as set forth above. Cells then are washed and incubated with a constant saturating concentration of a ¹²⁵I-mouse anti-human IgG in binding medium, with gentle agitation for 1 hour at 37°C. After extensive
5 washing, cells are released *via* trypsinization.

The mouse anti-human IgG employed above is directed against the Fc region of human IgG and can be obtained from Jackson ImmunoResearch Laboratories, Inc., West Grove, PA. The antibody is radioiodinated using the standard chloramine-T method. The antibody will bind to the Fc portion of any polypeptide/Fc protein that has bound to the cells. In all assays, non-
10 specific binding of ¹²⁵I-antibody is assayed in the absence of the Fc fusion protein/Fc, as well as in the presence of the Fc fusion protein and a 200-fold molar excess of unlabeled mouse anti-human IgG antibody.

Cell-bound ¹²⁵I-antibody is quantified on a Packard Autogamma counter. Affinity calculations (Scatchard, *Ann. N.Y. Acad. Sci.* 51:660, 1949) are generated on RS/1 (BBN
15 Software, Boston, MA) run on a Microvax computer.

Another type of suitable binding assay is a competitive binding assay. To illustrate, biological activity of a variant may be determined by assaying for the variant's ability to compete with the native protein for binding to the binding partner.

Competitive binding assays can be performed by conventional methodology. Reagents
20 that may be employed in competitive binding assays include radiolabeled IMX polypeptides and intact cells expressing the IMX polypeptide (endogenous or recombinant) on the cell surface. For example, a radiolabeled soluble IMX polypeptide fragment can be used to compete with a soluble IMX polypeptide variant for binding to the cell surface binding partner. Instead of intact cells, one could substitute a soluble binding partner/Fc fusion protein bound to a solid phase
25 through the interaction of Protein A or Protein G (on the solid phase) with the Fc moiety. Chromatography columns that contain Protein A and Protein G include those available from Pharmacia Biotech, Inc., Piscataway, NJ.

Another type of competitive binding assay utilizes the radiolabeled soluble binding partner, such as a soluble binding partner/Fc fusion protein, and intact cells expressing the IMX
30 polypeptide. Qualitative results can be obtained by competitive autoradiographic plate binding assays, while Scatchard plots (Scatchard, *Ann. N.Y. Acad. Sci.* 51:660, 1949) may be utilized to generate quantitative results.

USE OF IMX NUCLEIC ACIDS OR OLIGONUCLEOTIDES

In addition to being used to express polypeptides as described above, the nucleic acids of the invention, including DNA, and oligonucleotides thereof can be used:

- 5 — as probes to identify nucleic acid encoding proteins homologous to IMX polypeptides;
- to identify human chromosomes;
- to map genes on human chromosome numbers 7, 19, and 22;
- to identify genes associated with certain diseases, syndromes, or other conditions associated with human chromosome numbers 7, 19, and 22;
- 10 — as single-stranded sense or antisense oligonucleotides, to inhibit expression of polypeptide encoded by the IMX sequences;
- to help detect defective genes in an individual; and
- for gene therapy.

Probes

15 Among the uses of nucleic acids of the invention is the use of fragments as probes or primers. Such fragments generally comprise at least about 17 contiguous nucleotides of a DNA sequence. In other embodiments, a DNA fragment comprises at least 30, or at least 60, contiguous nucleotides of a DNA sequence.

20 Because homologs of SEQ ID NOs:1-26, from other mammalian species, are contemplated herein, probes based on the human DNA sequence of SEQ ID NOs:1-26 may be used to screen cDNA libraries derived from other mammalian species, using conventional cross-species hybridization techniques.

25 Using knowledge of the genetic code in combination with the amino acid sequences set forth above, sets of degenerate oligonucleotides can be prepared. Such oligonucleotides are useful as primers, e.g., in polymerase chain reactions (PCR), whereby DNA fragments are isolated and amplified.

Identifying Chromosome Number

All or a portion of the nucleic acids of SEQ ID NOs:1-26, including oligonucleotides, can be used by those skilled in the art using well-known techniques to identify the human chromosomes, and the specific locus thereof, that contain the DNA of IMX family members. Useful techniques include, but are not limited to, using the sequence or portions, including oligonucleotides, as a probe in various well-known techniques such as in situ hybridization to chromosome spreads, Southern blot hybridization to hybrid cell lines, fluorescent tagging, and radiation hybrid mapping.

For example, chromosomes can be mapped by radiation hybridization. PCR is performed using the Whitehead Institute/MIT Center for Genome Research Genebridge4 panel of 93 radiation hybrids (http://www-genome.wi.mit.edu/ftp/distribution/human_STS_releases/july97/rhmap/genebridge4.html). Primers are used which lie within a putative exon of the gene of interest and which amplify a product from human genomic DNA, but do not amplify hamster genomic DNA. The results of the PCRs are converted into a data vector that is submitted to the Whitehead/MIT Radiation Mapping site on the internet (<http://www-seq.wi.mit.edu>). The data is scored and the chromosomal assignment and placement relative to known Sequence Tag Site (STS) markers on the radiation hybrid map is provided. The following web site provides additional information about radiation hybrid mapping: http://www-genome.wi.mit.edu/ftp/distribution/human_STS_releases/july97/07-97.INTRO.html.

Identifying Associated Diseases

As described previously, IMX molecules numbered 4, 21, 44, and 56 have been mapped to particular chromosome locations. Thus, the nucleic acid of a particular IMX molecule or a fragment thereof can be used by one skilled in the art using well-known techniques to analyze abnormalities associated with gene mapping to such chromosomes. This enables one to distinguish conditions in which this marker is rearranged or deleted. In addition, nucleotides of such IMX molecules or fragments thereof can be used as a positional marker to map other genes of previously unknown location.

The DNA may be used in developing treatments for any disorder mediated (directly or indirectly) by defective, or insufficient amounts of, the genes corresponding to the nucleic acids of the invention. Disclosure herein of native nucleotide sequences permits the detection of defective genes, and the replacement thereof with normal genes. Defective genes may be

detected in *in vitro* diagnostic assays, and by comparison of a native nucleotide sequence disclosed herein with that of a gene derived from a person suspected of harboring a defect in this gene.

5 Sense-Antisense

Other useful fragments of the nucleic acids include antisense or sense oligonucleotides comprising a single-stranded nucleic acid sequence (either RNA or DNA) capable of binding to target mRNA (sense) or DNA (antisense) sequences. Antisense or sense oligonucleotides according to the present invention comprise a fragment of DNA (SEQ ID NOs:1-26). Such a
10 fragment generally comprises at least about 14 nucleotides, preferably from about 14 to about 30 nucleotides. The ability to derive an antisense or a sense oligonucleotide, based upon a cDNA sequence encoding a given protein is described in, for example, Stein and Cohen (*Cancer Res.* 48:2659, 1988) and van der Krol et al. (*BioTechniques* 6:958, 1988).

Binding of antisense or sense oligonucleotides to target nucleic acid sequences results in
15 the formation of duplexes that block or inhibit protein expression by one of several means, including enhanced degradation of the mRNA by RNaseH, inhibition of splicing, premature termination of transcription or translation, or by other means. The antisense oligonucleotides thus may be used to block expression of proteins. Antisense or sense oligonucleotides further comprise oligonucleotides having modified sugar-phosphodiester backbones (or other sugar
20 linkages, such as those described in WO91/06629) and wherein such sugar linkages are resistant to endogenous nucleases. Such oligonucleotides with resistant sugar linkages are stable *in vivo* (*i.e.*, capable of resisting enzymatic degradation) but retain sequence specificity to be able to bind to target nucleotide sequences.

Other examples of sense or antisense oligonucleotides include those oligonucleotides
25 which are covalently linked to organic moieties, such as those described in WO 90/10448, and other moieties that increases affinity of the oligonucleotide for a target nucleic acid sequence, such as poly-(L-lysine). Further still, intercalating agents, such as ellipticine, and alkylating agents or metal complexes may be attached to sense or antisense oligonucleotides to modify binding specificities of the antisense or sense oligonucleotide for the target nucleotide sequence.

30 Antisense or sense oligonucleotides may be introduced into a cell containing the target nucleic acid sequence by any gene transfer method, including, for example, lipofection, CaPO₄-mediated DNA transfection, electroporation, or by using gene transfer vectors such as Epstein-Barr virus.

Sense or antisense oligonucleotides also may be introduced into a cell containing the target nucleotide sequence by formation of a conjugate with a ligand binding molecule, as described in WO 91/04753. Suitable ligand binding molecules include, but are not limited to, cell surface receptors, growth factors, other cytokines, or other ligands that bind to cell surface receptors. Preferably, conjugation of the ligand binding molecule does not substantially interfere with the ability of the ligand binding molecule to bind to its corresponding molecule or receptor, or block entry of the sense or antisense oligonucleotide or its conjugated version into the cell.

Alternatively, a sense or an antisense oligonucleotide may be introduced into a cell containing the target nucleic acid sequence by formation of an oligonucleotide-lipid complex, as described in WO 90/10448. The sense or antisense oligonucleotide-lipid complex is preferably dissociated within the cell by an endogenous lipase.

USE OF IMX POLYPEPTIDES AND FRAGMENTED POLYPEPTIDES

Uses include, but are not limited to, the following:

- Purifying proteins and measuring activity thereof
- Delivery Agents
- Therapeutic and Research Reagents
- Molecular weight and Isoelectric focusing markers
- Controls for peptide fragmentation
- Identification of unknown proteins
- Preparation of Antibodies

Purification Reagents

Each of the polypeptides of the invention finds use as a protein purification reagent. The polypeptides may be attached to a solid support material and used to purify (binding partner) proteins by affinity chromatography. In particular embodiments, a polypeptide (in any form described herein that is capable of binding (binding partner)) is attached to a solid support by conventional procedures. As one example, chromatography columns containing functional groups that will react with functional groups on amino acid side chains of proteins are available (Pharmacia Biotech, Inc., Piscataway, NJ). In an alternative, a polypeptide/Fc protein (as discussed above) is attached to Protein A- or Protein G-containing chromatography columns through interaction with the Fc moiety.

The polypeptide also finds use in purifying or identifying cells that express (binding partner) on the cell surface. Polypeptides are bound to a solid phase such as a column chromatography matrix or a similar suitable substrate. For example, magnetic microspheres can be coated with the polypeptides and held in an incubation vessel through a magnetic field.

5 Suspensions of cell mixtures containing (binding partner) expressing cells are contacted with the solid phase having the polypeptides thereon. Cells expressing (binding partner) on the cell surface bind to the fixed polypeptides, and unbound cells then are washed away.

Alternatively, the polypeptides can be conjugated to a detectable moiety, then incubated with cells to be tested for (binding partner) expression. After incubation, unbound labeled
10 matter is removed and the presence or absence of the detectable moiety on the cells is determined.

In a further alternative, mixtures of cells suspected of containing (binding partner) cells are incubated with biotinylated polypeptides. Incubation periods are typically at least one hour in duration to ensure sufficient binding. The resulting mixture then is passed through a column
15 packed with avidin-coated beads, whereby the high affinity of biotin for avidin provides binding of the desired cells to the beads. Procedures for using avidin-coated beads are known (see Berenson, et al. *J. Cell. Biochem.*, 10D:239, 1986). Washing to remove unbound material, and the release of the bound cells, are performed using conventional methods.

20 Measuring Activity

Polypeptides also find use in measuring the biological activity of (binding partner) protein in terms of their binding affinity. The polypeptides thus may be employed by those conducting "quality assurance" studies, e.g., to monitor shelf life and stability of protein under different conditions. For example, the polypeptides may be employed in a binding affinity study
25 to measure the biological activity of a (binding partner) protein that has been stored at different temperatures, or produced in different cell types. The proteins also may be used to determine whether biological activity is retained after modification of a (binding partner) protein (e.g., chemical modification, truncation, mutation, etc.). The binding affinity of the modified (binding partner) protein is compared to that of an unmodified (binding partner) protein to detect any
30 adverse impact of the modifications on biological activity of (binding partner). The biological activity of a (binding partner) protein thus can be ascertained before it is used in a research study, for example.

Delivery Agents

The polypeptides also find use as carriers for delivering agents attached thereto to cells bearing binding partner. Cells expressing (binding partner) include those identified in (add
5 citation if reference known). The polypeptides thus can be used to deliver diagnostic or therapeutic agents to such cells (or to other cell types found to express (binding partner) on the cell surface) in *in vitro* or *in vivo* procedures.

Detectable (diagnostic) and therapeutic agents that may be attached to a polypeptide include, but are not limited to, toxins, other cytotoxic agents, drugs, radionuclides,
10 chromophores, enzymes that catalyze a colorimetric or fluorometric reaction, and the like, with the particular agent being chosen according to the intended application. Among the toxins are ricin, abrin, diphtheria toxin, *Pseudomonas aeruginosa* exotoxin A, ribosomal inactivating proteins, mycotoxins such as trichothecenes, and derivatives and fragments (e.g., single chains) thereof. Radionuclides suitable for diagnostic use include, but are not limited to, ^{123}I , ^{131}I , $^{99\text{m}}\text{Tc}$,
15 ^{111}In , and ^{76}Br . Examples of radionuclides suitable for therapeutic use are ^{131}I , ^{211}At , ^{77}Br , ^{186}Re , ^{188}Re , ^{212}Pb , ^{212}Bi , ^{109}Pd , ^{64}Cu , and ^{67}Cu .

Such agents may be attached to the polypeptide by any suitable conventional procedure. The polypeptide comprises functional groups on amino acid side chains that can be reacted with functional groups on a desired agent to form covalent bonds, for example. Alternatively, the
20 protein or agent may be derivatized to generate or attach a desired reactive functional group. The derivatization may involve attachment of one of the bifunctional coupling reagents available for attaching various molecules to proteins (Pierce Chemical Company, Rockford, Illinois). A number of techniques for radiolabeling proteins are known. Radionuclide metals may be attached to polypeptides by using a suitable bifunctional chelating agent, for example.

25 Conjugates comprising polypeptides and a suitable diagnostic or therapeutic agent (preferably covalently linked) are thus prepared. The conjugates are administered or otherwise employed in an amount appropriate for the particular application.

Therapeutic Agents

30 Polypeptides of the invention may be used in developing treatments for any disorder mediated (directly or indirectly) by defective, or insufficient amounts of the polypeptides. These polypeptides may be administered to a mammal afflicted with such a disorder.

The polypeptides may also be employed in inhibiting a biological activity of (binding partner), in *in vitro* or *in vivo* procedures. For example, a purified polypeptide may be used to

inhibit binding of (binding partner) to endogenous cell surface (binding partner). Biological effects that result from the binding of (binding partner) to endogenous receptors thus are inhibited.

IMX polypeptides may be administered to a mammal to treat a (binding partner)-mediated disorder. Such (binding partner)-mediated disorders include conditions caused (directly or indirectly) or exacerbated by (binding partner).

Compositions of the present invention may contain a polypeptide in any form described herein, such as native proteins, variants, derivatives, oligomers, and biologically active fragments. In particular embodiments, the composition comprises a soluble polypeptide or an oligomer comprising soluble polypeptides.

Compositions comprising an effective amount of a polypeptide of the present invention, in combination with other components such as a physiologically acceptable diluent, carrier, or excipient, are provided herein. The polypeptides can be formulated according to known methods used to prepare pharmaceutically useful compositions. They can be combined in admixture, either as the sole active material or with other known active materials suitable for a given indication, with pharmaceutically acceptable diluents (e.g., saline, Tris-HCl, acetate, and phosphate buffered solutions), preservatives (e.g., thimerosal, benzyl alcohol, parabens), emulsifiers, solubilizers, adjuvants and/or carriers. Suitable formulations for pharmaceutical compositions include those described in *Remington's Pharmaceutical Sciences*, 16th ed. 1980, Mack Publishing Company, Easton, PA.

In addition, such compositions can be complexed with polyethylene glycol (PEG), metal ions, or incorporated into polymeric compounds such as polyacetic acid, polyglycolic acid, hydrogels, dextran, etc., or incorporated into liposomes, microemulsions, micelles, unilamellar or multilamellar vesicles, erythrocyte ghosts or spheroblasts. Such compositions will influence the physical state, solubility, stability, rate of *in vivo* release, and rate of *in vivo* clearance, and are thus chosen according to the intended application.

The compositions of the invention can be administered in any suitable manner, e.g., topically, parenterally, or by inhalation. The term "parenteral" includes injection, e.g., by subcutaneous, intravenous, or intramuscular routes, also including localized administration, e.g., at a site of disease or injury. Sustained release from implants is also contemplated. One skilled in the pertinent art will recognize that suitable dosages will vary, depending upon such factors as the nature of the disorder to be treated, the patient's body weight, age, and general condition, and the route of administration. Preliminary doses can be determined according to animal tests, and

the scaling of dosages for human administration is performed according to art-accepted practices.

Compositions comprising nucleic acids in physiologically acceptable formulations are also contemplated. DNA may be formulated for injection, for example.

5

Research Agents

Another use of the polypeptide of the present invention is as a research tool for studying the biological effects that result from inhibiting interactions between polypeptides of the invention and their "binding partner". Polypeptides also may be employed in *in vitro* assays for
10 detecting binding partner or cells expressing the binding partner or the interactions thereof.

Molecular Weight, Isoelectric Point Markers

The polypeptides of the present invention can be subjected to fragmentation into smaller peptides by chemical and enzymatic means, and the peptide fragments so produced can be used in the analysis of other proteins or polypeptides. For example, such peptide fragments can be
15 used as peptide molecular weight markers, peptide isoelectric point markers, or in the analysis of the degree of peptide fragmentation. Thus, the invention also includes these polypeptides and peptide fragments, as well as kits to aid in the determination of the apparent molecular weight and isoelectric point of an unknown protein and kits to assess the degree of fragmentation of an unknown protein.

20 Although all methods of fragmentation are encompassed by the invention, chemical fragmentation is a preferred embodiment, and includes the use of cyanogen bromide to cleave under neutral or acidic conditions such that specific cleavage occurs at methionine residues (E. Gross, *Methods in Enz.* 11:238-255, 1967). This can further include additional steps, such as a carboxymethylation step to convert cysteine residues to an unreactive species.

25 Enzymatic fragmentation is another preferred embodiment, and includes the use of a protease such as Asparaginylendo-peptidase, Arginylendo-peptidase, *Achromobacter* protease I, Trypsin, *Staphylococcus aureus* V8 protease, Endoproteinase Asp-N, or Endoproteinase Lys-C under conventional conditions to result in cleavage at specific amino acid residues.

Asparaginylendo-peptidase can cleave specifically on the carboxyl side of the asparagine
30 residues present within the polypeptides of the invention. Arginylendo-peptidase can cleave specifically on the carboxyl side of the arginine residues present within these polypeptides. *Achromobacter* protease I can cleave specifically on the carboxyl side of the lysine residues present within the polypeptides (Sakiyama and Nakat, U.S. Patent No. 5,248,599; T. Masaki et al., *Biochim. Biophys. Acta* 660:44-50, 1981; T. Masaki et al., *Biochim. Biophys. Acta* 660:51-

55, 1981). Trypsin can cleave specifically on the carboxyl side of the arginine and lysine residues present within polypeptides of the invention. Enzymatic fragmentation may also occur with a protease that cleaves at multiple amino acid residues. For example, *Staphylococcus aureus* V8 protease can cleave specifically on the carboxyl side of the aspartic and glutamic acid residues present within polypeptides (D. W. Cleveland, *J. Biol. Chem.* 3:1102-1106, 1977). Endoproteinase Asp-N can cleave specifically on the amino side of the asparagine residues present within polypeptides. Endoproteinase Lys-C can cleave specifically on the carboxyl side of the lysine residues present within polypeptides of the invention. Other enzymatic and chemical treatments can likewise be used to specifically fragment these polypeptides into a unique set of specific peptides.

Of course, the peptides and fragments of the polypeptides of the invention can also be produced by conventional recombinant processes and synthetic processes well known in the art. With regard to recombinant processes, the polypeptides and peptide fragments encompassed by invention can have variable molecular weights, depending upon the host cell in which they are expressed. Glycosylation of polypeptides and peptide fragments of the invention in various cell types can result in variations of the molecular weight of these pieces, depending upon the extent of modification. The size of these pieces can be most heterogeneous with fragments of polypeptide derived from the extracellular portion of the polypeptide. Consistent polypeptides and peptide fragments can be obtained by using polypeptides derived entirely from the transmembrane and cytoplasmic regions, pretreating with N-glycanase to remove glycosylation, or expressing the polypeptides in bacterial hosts.

The molecular weight of these polypeptides can also be varied by fusing additional peptide sequences to both the amino and carboxyl terminal ends of polypeptides of the invention. Fusions of additional peptide sequences at the amino and carboxyl terminal ends of polypeptides of the invention can be used to enhance expression of these polypeptides or aid in the purification of the protein. In addition, fusions of additional peptide sequences at the amino and carboxyl terminal ends of polypeptides of the invention will alter some, but usually not all, of the fragmented peptides of the polypeptides generated by enzymatic or chemical treatment. Of course, mutations can be introduced into polypeptides of the invention using routine and known techniques of molecular biology. For example, a mutation can be designed so as to eliminate a site of proteolytic cleavage by a specific enzyme or a site of cleavage by a specific chemically induced fragmentation procedure. The elimination of the site will alter the peptide fingerprint of polypeptides of the invention upon fragmentation with the specific enzyme or chemical procedure.

The polypeptides and the resultant fragmented peptides can be analyzed by methods including sedimentation, electrophoresis, chromatography, and mass spectrometry to determine their molecular weights. Because the unique amino acid sequence of each piece specifies a molecular weight, these pieces can thereafter serve as molecular weight markers using such analysis techniques to assist in the determination of the molecular weight of an unknown protein, polypeptides or fragments thereof. The molecular weight markers of the invention serve particularly well as molecular weight markers for the estimation of the apparent molecular weight of proteins that have similar apparent molecular weights and, consequently, allow increased accuracy in the determination of apparent molecular weight of proteins.

When the invention relates to the use of fragmented peptide molecular weight markers, those markers are preferably at least 10 amino acids in size. More preferably, these fragmented peptide molecular weight markers are between 10 and 100 amino acids in size. Even more preferable are fragmented peptide molecular weight markers between 10 and 50 amino acids in size and especially between 10 and 35 amino acids in size. Most preferable are fragmented peptide molecular weight markers between 10 and 20 amino acids in size.

Among the methods for determining molecular weight are sedimentation, gel electrophoresis, chromatography, and mass spectrometry. A particularly preferred embodiment is denaturing polyacrylamide gel electrophoresis (U. K. Laemmli, *Nature* 227:680-685, 1970). Conventionally, the method uses two separate lanes of a gel containing sodium dodecyl sulfate and a concentration of acrylamide between 6-20%. The ability to simultaneously resolve the marker and the sample under identical conditions allows for increased accuracy. It is understood, of course, that many different techniques can be used for the determination of the molecular weight of an unknown protein using polypeptides of the invention, and that this embodiment in no way limits the scope of the invention.

Each unglycosylated polypeptide or fragment thereof has a pI that is intrinsically determined by its unique amino acid sequence (which pI can be estimated by the skilled artisan using any of the computer programs designed to predict pI values currently available, calculated using any well-known amino acid pKa table, or measured empirically). Therefore these polypeptides and fragments thereof can serve as specific markers to assist in the determination of the isoelectric point of an unknown protein, polypeptide, or fragmented peptide using techniques such as isoelectric focusing. These polypeptide or fragmented peptide markers serve particularly well for the estimation of apparent isoelectric points of unknown proteins that have apparent isoelectric points close to that of the polypeptide or fragmented peptide markers of the invention.

The technique of isoelectric focusing can be further combined with other techniques such as gel electrophoresis to simultaneously separate a protein on the basis of molecular weight and charge. The ability to simultaneously resolve these polypeptide or fragmented peptide markers and the unknown protein under identical conditions allows for increased accuracy in the determination of the apparent isoelectric point of the unknown protein. This is of particular interest in techniques, such as two dimensional electrophoresis (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)), where the nature of the procedure dictates that any markers should be resolved simultaneously with the unknown protein. In addition, with such methods, these polypeptides and fragmented peptides thereof can assist in the determination of both the isoelectric point and molecular weight of an unknown protein or fragmented peptide.

Polypeptides and fragmented peptides can be visualized using two different methods that allow a discrimination between the unknown protein and the molecular weight markers. In one embodiment, the polypeptide and fragmented peptide molecular weight markers of the invention can be visualized using antibodies generated against these markers and conventional immunoblotting techniques. This detection is performed under conventional conditions that do not result in the detection of the unknown protein. It is understood that it may not be possible to generate antibodies against all polypeptide fragments of the invention, since small peptides may not contain immunogenic epitopes. It is further understood that not all antibodies will work in this assay; however, those antibodies which are able to bind polypeptides and fragments of the invention can be readily determined using conventional techniques.

The unknown protein is also visualized by using a conventional staining procedure. The molar excess of unknown protein to polypeptide or fragmented peptide molecular weight markers of the invention is such that the conventional staining procedure predominantly detects the unknown protein. The level of these polypeptide or fragmented peptide molecular weight markers is such as to allow little or no detection of these markers by the conventional staining method. The preferred molar excess of unknown protein to polypeptide molecular weight markers of the invention is between 2 and 100,000 fold. More preferably, the preferred molar excess of unknown protein to these polypeptide molecular weight markers is between 10 and 10,000 fold and especially between 100 and 1,000 fold.

It is understood of course that many techniques can be used for the determination and detection of molecular weight and isoelectric point of an unknown protein, polypeptides, and fragmented peptides thereof using these polypeptide molecular weight markers and peptide fragments thereof and that these embodiments in no way limit the scope of the invention.

In another embodiment, the analysis of the progressive fragmentation of the polypeptides of the invention into specific peptides (D. W. Cleveland et al., *J. Biol. Chem.* 252:1102-1106, 1977), such as by altering the time or temperature of the fragmentation reaction, can be used as a control for the extent of cleavage of an unknown protein. For example, cleavage of the same amount of polypeptide and unknown protein under identical conditions can allow for a direct comparison of the extent of fragmentation. Conditions that result in the complete fragmentation of the polypeptide can also result in complete fragmentation of the unknown protein.

As to the specific use of the polypeptides and fragmented peptides of the invention as molecular weight markers, the fragmentation of the polypeptides of SEQ ID NOs:27-38 with cyanogen bromide generates a unique set of fragmented peptide molecular weight markers. An additional fragment results if the initiating methionine is present. The distribution of methionine residues determines the number of amino acids in each peptide and the unique amino acid composition of each peptide determines its molecular weight.

In addition, the preferred purified polypeptides of the invention (SEQ ID NOs:27-38) have calculated molecular weights of approximately 3683, 1783, 11248, 75503, 43040, 8051, 33306, 3515, 10736, 25162, and 2450 Daltons.

Where an intact protein is used, the use of these polypeptide molecular weight markers allows increased accuracy in the determination of apparent molecular weight of proteins that have apparent molecular weights close to 3683, 1783, 11248, 75503, 43040, 8051, 33306, 3515, 10736, 25162, or 2450 Daltons. Where fragments are used, there is increased accuracy in determining molecular weight over the range of the molecular weights of the fragment.

Finally, as to the kits that are encompassed by the invention, the constituents of such kits can be varied, but typically contain the polypeptide and fragmented peptide molecular weight markers. Also, such kits can contain the polypeptides wherein a site necessary for fragmentation has been removed. Furthermore, the kits can contain reagents for the specific cleavage of the polypeptide and the unknown protein by chemical or enzymatic cleavage. Kits can further contain antibodies directed against polypeptides or fragments thereof of the invention.

Identification of Unknown Proteins

As set forth above, a polypeptide or peptide fingerprint can be entered into or compared to a database of known proteins to assist in the identification of the unknown protein using mass spectrometry (W.J. Henzel et al., *Proc. Natl. Acad. Sci. USA* 90:5011-5015, 1993; D. Fenyo et al., *Electrophoresis* 19:998-1005, 1998). A variety of computer software programs to facilitate these comparisons are accessible via the Internet, such as Protein Prospector (Internet site:

prospector.uscf.edu). MultiIdent (Internet site: www.expasy.ch/sprot/multiident.html), PeptideSearch (Internet site: www.mann.embl-heidelberg.de...deSearch/FR_PeptideSearchForm.html), and ProFound (Internet site: www.chait-sgi.rockefeller.edu/cgi-bin/prot-id-frag.html). These programs allow the user to specify the cleavage agent and the molecular weights of the fragmented peptides within a designated tolerance. The programs compare these molecular weights to protein databases to assist in determining the identity of the unknown protein.

In addition, a polypeptide or peptide digest can be sequenced using tandem mass spectrometry (MS/MS) and the resulting sequence searched against databases (J.K. Eng, et al., *J. Am. Soc. Mass Spec.* 5:976-989 (1994); M. Mann and M. Wilm, *Anal. Chem.* 66:4390-4399 (1994); J.A. Taylor and R.S. Johnson, *Rapid Comm. Mass Spec.* 11:1067-1075 (1997)). Searching programs that can be used in this process exist on the Internet, such as Lutefisk 97 (Internet site: www.lsbcc.com:70/Lutefisk97.html), and the Protein Prospector, Peptide Search and ProFound programs described above. Therefore, adding the sequence of a gene and its predicted protein sequence and peptide fragments to a sequence database can aid in the identification of unknown proteins using tandem mass spectrometry.

Antibodies

Antibodies that are immunoreactive with the polypeptides of the invention are provided herein. Such antibodies specifically bind to the polypeptides *via* the antigen-binding sites of the antibody (as opposed to non-specific binding). Thus, the polypeptides, fragments, variants, fusion proteins, etc., as set forth above may be employed as immunogens in producing antibodies immunoreactive therewith.

Polyclonal and monoclonal antibodies may be prepared by conventional techniques. See, for example, *Monoclonal Antibodies, Hybridomas: A New Dimension in Biological Analyses*, Kennet et al. (eds.), Plenum Press, New York (1980); and *Antibodies: A Laboratory Manual*, Harlow and Land (eds.), Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, (1988).

Antigen-binding fragments of such antibodies, which may be produced by conventional techniques, are also encompassed by the present invention. Examples of such fragments include, but are not limited to, Fab and F(ab')₂ fragments. Antibody fragments and derivatives produced by genetic engineering techniques are also provided.

The monoclonal antibodies of the present invention include chimeric antibodies, e.g., humanized versions of murine monoclonal antibodies. Such humanized antibodies may be

prepared by known techniques, and offer the advantage of reduced immunogenicity when the antibodies are administered to humans. In one embodiment, a humanized monoclonal antibody comprises the variable region of a murine antibody (or just the antigen binding site thereof) and a constant region derived from a human antibody. Alternatively, a humanized antibody

5 fragment may comprise the antigen binding site of a murine monoclonal antibody and a variable region fragment (lacking the antigen-binding site) derived from a human antibody. Procedures for the production of chimeric and further engineered monoclonal antibodies include those described in Riechmann et al. (*Nature* 332:323, 1988), Liu et al. (*PNAS* 84:3439, 1987), Larrick et al. (*Bio/Technology* 7:934, 1989), and Winter and Harris (*TIPS* 14:139, May, 1993).

10 Procedures to generate antibodies transgenically can be found in GB 2,272,440, US Patent Nos. 5,569,825 and 5,545,806 and related patents claiming priority therefrom, all of which are incorporated by reference herein.

In one embodiment, the antibodies are specific for the polypeptides of the present invention, and do not cross-react with other proteins. Screening procedures by which such

15 antibodies may be identified are well known, and may involve immunoaffinity chromatography, for example.

Hybridoma cell lines that produce monoclonal antibodies specific for the polypeptides of the invention are also contemplated herein. Such hybridomas may be produced and identified by conventional techniques. One method for producing such a hybridoma cell line comprises

20 immunizing an animal with a polypeptide; harvesting spleen cells from the immunized animal; fusing said spleen cells to a myeloma cell line, thereby generating hybridoma cells; and identifying a hybridoma cell line that produces a monoclonal antibody that binds the polypeptide. The monoclonal antibodies may be recovered by conventional techniques.

25 Uses Thereof

The antibodies of the invention can be used in assays to detect the presence of the polypeptides or fragments of the invention, either *in vitro* or *in vivo*. The antibodies also may be employed in purifying polypeptides or fragments of the invention by immunoaffinity chromatography.

30 Those antibodies that additionally can block binding of the polypeptides of the invention to the binding partner may be used to inhibit a biological activity that results from such binding. Such blocking antibodies may be identified using any suitable assay procedure, such as by testing antibodies for the ability to inhibit binding of the binding partner to certain cells expressing the binding partner. Alternatively, blocking antibodies may be identified in assays

for the ability to inhibit a biological effect that results from binding of the binding partner to target cells.

Such an antibody may be employed in an *in vitro* procedure, or administered *in vivo* to inhibit a biological activity mediated by the entity that generated the antibody. Disorders caused or exacerbated (directly or indirectly) by the interaction of (binding partner) with cell surface (binding partner) receptor thus may be treated. A therapeutic method involves *in vivo* administration of a blocking antibody to a mammal in an amount effective in inhibiting a (binding partner)-mediated biological activity. Monoclonal antibodies are generally preferred for use in such therapeutic methods. In one embodiment, an antigen-binding antibody fragment is employed.

Antibodies may be screened for agonistic (*i.e.*, ligand-mimicking) properties. Such antibodies, upon binding to cell surface antigen, induce biological effects (e.g., transduction of biological signals) similar to the biological effects induced when (binding partner) binds to cell surface antigen.

Compositions comprising an antibody that is directed against a polypeptide of the invention, and a physiologically acceptable diluent, excipient, or carrier, are provided herein.

Also provided herein are conjugates comprising a detectable (e.g., diagnostic) or therapeutic agent, attached to the antibody. Examples of such agents are presented above. The conjugates find use in *in vitro* or *in vivo* procedures.

The following examples are provided to further illustrate particular embodiments of the invention, and are not to be construed as limiting the scope of the present invention.

EXAMPLE 1: Isolation of the IMX Nucleic Acids

The T84 Epithelial Barrier Model

As discussed above, damage to the intestinal epithelial barrier is a hallmark of (IBD), and a number of *in vitro* models of epithelial barrier function have been developed over the years. The best characterized of these models is the T84 intestinal epithelial barrier system, Dharmasathaphorn et al., *Am. J. Physiol.*, 246:G204-G208, 1984 and Madara et al., *J. Cell Biol.*, 101:2124-2133, 1985).

T84 cells were plated on 75 mm polycarbonate transwell filter inserts (Costar) and grown in DME/F12 (1:1) containing 10% heat-inactivated bovine calf serum. The cells were maintained at confluence for 2-3 days, and integrity of the epithelial barrier was determined by measuring transepithelial electrical resistance (TER) using an EVOM epithelial voltohmmeter (World Precision Instruments). When the TER values were greater than 1000 ohms/cm² and

were stable, cells were treated with interferon-g (30 ng/ml. Genzyme) added to the basolateral side of the membrane. At various times after treatment (4, 24 and 44h), TERs were measured to monitor the interferon-induced disruption of the barrier, and RNA was harvested from the cells at those time points using TRIzol reagent (Life Technologies). RNA was extracted using conventional methods and subsequently used for TOGA™ analysis as described in Example 2.

TOGA™ analysis

This example describes a method for determining mRNA expression characteristics. The isolated RNA was analyzed using a method of simultaneous sequence-specific identification of mRNAs known as TOGA™ (Total Gene expression Analysis) described in U.S. Patent No. 5,459,037 and U.S. Patent No. 5,807,680; hereby incorporated herein by reference. Preferably, prior to the application of the TOGA™ technique, the isolated RNA was enriched to form a starting polyA-containing mRNA population by methods known in the art. In a preferred embodiment, the TOGA™ method further comprised an additional PCR step performed using four separate reactions, one for each of the four 5' PCR primers, and cDNA templates prepared from a population of antisense cRNAs. A final PCR step used 256 5' PCR primers in 64 subpools for each of the four reactions of the previous step produced PCR products that were cDNA fragments that corresponded to the 3'-region of the starting mRNA population.

The produced PCR products were then identified by a) the sequence of at least the 5' seven base pairs, preferably the sequence of the entire fragment, and b) the length of the fragment. These two parameters, sequence and fragment length, were used to compare the obtained PCR products to a database of known polynucleotide sequences. A database search for homologous sequences in Genbank resulted in no matches, indicating the novelty of the IMX sequences of the invention. The intensities of the PCR products were compared across the 4 input RNA samples (t=0, 4hr, 24hr, 44hr) and species that were regulated were identified and further characterized (Table 1).

The DSTs of SEQ ID NO:1-26 and fragments thereof are useful as probes to study and diagnose the changes in gene expression demonstrated by the data of Table 1. Alternatively, polypeptides and fragments thereof that are the translations of SEQ ID NO:1-26 and fragments thereof are useful as probes to study and diagnose the changes in gene expression demonstrated by the data of Table 1.

Name	DST SEQ ID NO:	Digital Address (<u>Msp</u> I)	Relative PCR Fragment Amount			
			0	4hr	24hr	44hr
IMX 4	1	TAAG 187	41	94	350	299
IMX 10	2	TTCT 159	250	716	250	207
IMX 21	3	CTTC 364	35	60	330	749
IMX 28	4	CCAT 414	138	435	647	464
IMX 32	5	CCGA 374	237	539	25	32
IMX 39	6	TGGA 450	24	87	110	78
IMX 40	7	TCTA 163	38	294	67	153
IMX 42	8	GAGC 426	145	733	876	1076
IMX 44	9	CTGC 221	248	344	1499	1624
IMX 56	10	GATA 111	955	1223	2357	3450

Table 1. IMX Nucleic Acid Molecules. Relative expression levels determined by TOGA™.

5

EXAMPLE 2: Further Characterization of IMX4

Nucleic Acid Molecules and Polypeptides

An additional 600bp sequence (SEQ ID NO:11) that included DST IMX4 (SEQ ID NO:1) was obtained by anchor PCR from a T84 library. A clone approximately 2.0kb in length was then isolated from a commercial library (Origene). The sequence of the 3' end of the clone (SEQ ID NO:12) included the sequence of the IMX4 DST. The sequence of the 5' end of the clone (SEQ ID NO:13) matched part of the human ApoL gene (AF019225, Figure 21). Complete sequence of the clone was not obtained. However, comparison of 5' end of the clone suggests it represents an alternative splice product to the reported ApoL sequence, i.e., bases 1-168 match to 2 exons on PAC carrying the ApoL gene but are not included in the reported complete cDNA. See Figure 22. A polypeptide translated from IMX4 is provided in SEQ ID NO: 27.

EXAMPLE 3: Further Characterization of IMX10 and IMX21

Nucleic Acid Molecules and Polypeptides

Further experiments failed to show regulation of these DSTs in replications using PCR with extended primers and Northern blots. A polypeptide translated from IMX10 (SEQ ID

NO:14) is provided in SEQ ID NO: 28. A polypeptide translated from IMX21 (SEQ ID NO:15) is provided in SEQ ID NO: 29.

EXAMPLE 4: Further Characterization of IMX28

Nucleic Acid Molecules and Polypeptides

One polypeptide translated from IMX28 is provided in SEQ ID NO: 30.

A GST fusion protein of IMX28 was used in studies designed to identify potential ligands for this enzyme. The GST fusion of IMX28 was prepared as follows: IMX28-8, a clone with the full length coding sequence, was excised from pGEM by SalI-NotI digestion and subcloned into SalI-NotI digested pGEX-5X-3 (Pharmacia). The pGEX-5X-3 plasmid contains the Schistosoma japonicum glutathione-S-transferase (GST) gene 5' to the SalI-NotI cloning site. Insertion of the IMX28-8 DNA fragment into this cloning site results in a fusion gene containing GST-IMX28 in the proper reading frame. The complete sequence of the fusion protein consists of GST followed by a Factor Xa tripeptide cleavage site, the peptide consisting of Gly-Ile-Pro-Arg-Asn-Ser-Arg-Val-Asp-Ala-Thr that is derived partially from the pGEX vector and partially from the SalI-NotI fragment of IMX28 and IMX28. The vector containing the fusion gene was electroporated into competent bacterial cells and a single colony containing the DNA, confirmed by diagnostic restriction digestion, was expanded to a 250ml culture for purification of the fusion protein.

Briefly, cells from these cultures were pelleted and resuspended in STE buffer containing a protease inhibitor cocktail and 20ug/ml hen egg lysozyme. The cells were lysed by addition of 10% sarcosyl in STE (final sarcosyl concentration was 1.5%), sonicated and centrifuged to remove cell debris. The supernatant was passed over a Glutathione-Sepharose 4B column, washed sequentially with 1.5% sarcosyl in STE and then STE. The bound material, containing the GST-IMX28 fusion protein was eluted by adding 20mM Glutathione in STE. The eluted material was analyzed by SDS-PAGE to confirm the presence of the GST-IMX28 fusion protein.

A recombinant adenovirus construct expressing IMX28 was used to infect T84 cells to determine effects on epithelial barrier formation. To construct adenoviral vectors with untagged full length Imx28 and NH2-terminally FLAG tagged Imx28, the pADEASY1 system described by He et al. (1998) Proc. Natl. Acad. Sci. USA 95: 2509-2514, was used. For the untagged construct a NotI fragment carrying the full coding region of Imx28 was inserted into the polylinker of the pAdtrackCMV shuttle vector. After confirmation of the correct orientation the

vector was cut with the restriction enzyme PacI and co-transformed into E. coli. Strain BJ5183 along with the pADEasy1 vector. Recombinants containing the Imx28 product were screened for as described and by gene specific PCR. To generate non-replicative, infectious viral particles the helper strain 293EBNA-EBNA or 293MSR was transfected with PmeI linearized recombinant vector. After 3 days virus was harvested by freeze-thaw lysis of the transfected cells. Several rounds of infection of these cells with viral supernatants were performed to boost titers. The tagged version was prepared by addition of the FLAG sequence and a Gly-Gly-Gly-Gly spacer region onto the full length coding sequence of Imx28 via PCR with a 5' oligo with the incorporated nucleotides for the tag and spacer. The PCR product was cloned into a TA- vector and excised with NotI and cloned as above. Screening and viral particle production was as above. No deleterious or positive effects were observed on T84 barrier function in the presence or absence of IFN-gamma.

EXAMPLE 5: Further Characterization of IMX32

Nucleic Acid Molecules and Polypeptides

The original clone produced by extending the IMX32 DST was 1588 bp in length (SEQ ID NO: 17). The first round of anchor PCR yielded a 670bp product that was used to produce a contig that extended the sequence 585 bases to 2173 (SEQ ID NO: 17). Another round of anchor PCR yielded the 1676 bp product, immunex32-a24.seq. (SEQ ID NO: 18) which further extended the contig for Imx32 to 2834 bp. In addition, the immunex32-a24.seq product revealed an alternative splice lacking bases 211-639 of IMX32-1 clone.

There is an ORF for 155 amino acids at the 5' end of the sequence (SEQ ID NO:31 translated from SEQ ID NO: 16; SEQ ID NO:32 translated from SEQ ID NO: 18). BLAST analysis suggests that the ORF may contain a KRAB-like domain because it has partial matches to Zn-finger proteins which contain KRAB domains at their N-terminus such as ZNF140.

EXAMPLE 6: Further Characterization of IMX39

Nucleic Acid Molecules and Polypeptides

One polypeptide translated from IMX39 is provided in SEQ ID NO: 33.

IMX39 FLAG-tagged and untagged adenoviral vector versions were prepared as outlined above for Imx28 in Example 4, above. Controls for the expression were infection with other FLAG- tagged adenoviral delivered proteins such as IMX5. Based upon the predicted structure from its cDNA sequence, it was expected that the IMX39 polypeptide would be a cytoplasmic protein. However, since some secreted cytokines, such as those of the IL-1 family, lack a

predicted signal sequence but are secreted from the cell, this expectation was tested experimentally in both T84 cells via adenoviral mediated transduction and via transfection into CV1EBNA cells with another expression vector, pDC412. Despite evidence for successful expression in the transfected cells, no product was found in the media as determined by FLAG western (T84 system) or 35-S radiolabeled product (CV1EBNA system). The positive control in the CV1 system was an Imx44-Fc fusion protein known to be secreted. In both systems empty vectors were used as a negative control. The attempted expression of IMX39 polypeptides had no apparent effect on barrier function in the absence or presence of IFN-gamma.

10

EXAMPLE 7: Further Characterization of IMX40

Nucleic Acid Molecules and Polypeptides

Anchor PCR using an Origene human small intestine library revealed a product of 1669 bp (SEQ ID NO: 20). Bases 1-265 of the DST IMX40 (SEQ ID NO:7) align with bases 1565-1669 of this sequence. The anchor PCR product has a potential 155 aa ORF from bases 185-650. A translation of this ORF is given in SEQ ID NO:34. This ORF has no recognizable features, motifs, or database homologies.

20

EXAMPLE 8: Further Characterization of IMX42

Nucleic Acid Molecules and Polypeptides

Several attempts to screen libraries using DST IMX42 (SEQ ID NO:8) as well as anchor PCR and RACE have not produced a clone with a longer sequence. BLAST comparison against imxhutdb extended sequence approx 200 bases to 592 bases (SEQ ID NO: 21). However, the sequence extension did not increase any database matches found by comparison to the DST IMX42 sequence. The contig has a potential ORF of 134 amino acids (SEQ ID NO: 36) which, while a COOH terminal extension of SEQ ID NO:35, may be a partial sequence.

EXAMPLE 9: Further Characterization of IMX44**Nucleic Acid Molecules and Polypeptides**

5 Longer polynucleotides corresponding to DST IMX44 have been identified (SEQ ID NO:22, 23, 24). A translated polypeptide is found in SEQ ID NO:37. A soluble Fc form of IMX44 polypeptide was synthesized and used in various assays. IMX44-Fc had no effect on T84 barrier function in the absence or presence of IFN-gamma. IMX44-Fc had no effect on natural killer (NK) cell activation. IMX44-Fc had no positive hits on cognate screen assays.

10 A soluble FLAG polyHis form was also produced. No activity in cellular activation assays was found nor any alteration of cytokine production using this polypeptide in assays.

The expression of IMX44 in various murine models of gut inflammation was determined by Northern and array analysis. Little to no regulation of transcript was found in anti-CD3-induced ileitis in C57BL/6 mice, DSS-induced colitis in BALB/c mice or C57BL/6
15 mice, *mdr1* knock out mice with colitis, and IFN-gamma stimulated LN T cells.

EXAMPLE 10: Further Characterization of IMX56**Nucleic Acid Molecules and Polypeptides**

A translated IMX56 polypeptide is found in SEQ ID NO:38. Comparison of the DST
20 sequence IMX56 to IMAGE consortium clones extended the sequence (SEQ ID NO:25) which was 3' on sequenced PAC to the described end of human ApoL. Anchor PCR using a T84 library produced results that indicated that the IMX56 DST is derived from an alternate 3' UTR of ApoL.

EXAMPLE 11: Monoclonal Antibodies That Bind IMX Polypeptides

25 Monoclonal antibodies that bind the polypeptides of the invention can be prepared by methods well known in the art. Suitable immunogens that may be employed in generating such antibodies include, but are not limited to, purified IMX polypeptides or an immunogenic fragment thereof such as the extracellular domain, or fusion proteins containing IMX
30 polypeptides (e.g., a soluble IMX 21 polypeptide/Fc fusion protein).

Purified IMX polypeptides of the invention can be used to generate monoclonal antibodies immunoreactive therewith, using conventional techniques such as those described in U.S. Patent 4,411,993. Briefly, mice are immunized with an IMX polypeptide immunogen emulsified in complete Freund's adjuvant, and injected in amounts ranging from 10-100 :g

subcutaneously or intraperitoneally. Ten to twelve days later, the immunized animals are boosted with additional IMX polypeptide immunogen emulsified in incomplete Freund's adjuvant. Mice are periodically boosted thereafter on a weekly to bi-weekly immunization schedule. Serum samples are periodically taken by retro-orbital bleeding or tail-tip excision to
5 test for anti-IMX polypeptide antibodies by dot blot assay, ELISA (Enzyme-Linked Immunosorbent Assay) or inhibition of IMX polypeptide/binding partner interactions.

Following detection of an appropriate antibody titer, positive animals are provided one last intravenous injection of IMX polypeptide immunogen in saline. Three to four days later, the animals are sacrificed, spleen cells harvested, and spleen cells are fused to a murine myeloma
10 cell line, e.g., NS1 or preferably P3x63Ag8.653 (ATCC CRL 1580). Fusions generate hybridoma cells, which are plated in multiple microtiter plates in a HAT (hypoxanthine, aminopterin and thymidine) selective medium to inhibit proliferation of non-fused cells, myeloma hybrids, and spleen cell hybrids.

The hybridoma cells are screened by ELISA for reactivity against the purified IMX
15 polypeptide of interest by adaptations of the techniques disclosed in Engvall et al., *Immunochem.* 8:871, 1971 and in U.S. Patent 4,703,004. A preferred screening technique is the antibody capture technique described in Beckmann et al., (*J. Immunol.* 144:4212, 1990). Positive hybridoma cells can be injected intraperitoneally into syngeneic BALB/c mice to produce ascites containing high concentrations of anti-IMX polypeptide monoclonal antibodies.
20 Alternatively, hybridoma cells can be grown *in vitro* in flasks or roller bottles by various techniques. Monoclonal antibodies produced in mouse ascites can be purified by ammonium sulfate precipitation, followed by gel exclusion chromatography. Alternatively, affinity chromatography based upon binding of antibody to Protein A or Protein G can also be used, as can affinity chromatography based upon binding to the IMX polypeptide of interest.

25 Alternatively, IMX nucleic acid molecules or fragments thereof can be expressed to produce IMX polypeptides or fragments thereof, that can be used to make antibodies that are useful for identifying corresponding polypeptides in techniques such as western blotting, immunocytochemistry, and ELISA assays using standard techniques such as those described in U.S. Patent No. 4,900,811, incorporated by reference herein.

30 The references cited herein are incorporated by reference herein in their entirety.

We claim:

1. A purified polypeptide selected from the group consisting of:

- a) a polypeptide comprising an amino acid sequence of SEQ ID NO:27;
- b) a polypeptide comprising an amino acid sequence of SEQ ID NO:28;
- 5 c) a polypeptide comprising an amino acid sequence of SEQ ID NO:29;
- d) a polypeptide comprising an amino acid sequence of SEQ ID NO:30;
- e) a polypeptide comprising an amino acid sequence of SEQ ID NO:31;
- f) a polypeptide comprising an amino acid sequence of SEQ ID NO:32;
- 10 g) a polypeptide comprising an amino acid sequence of SEQ ID NO:33;
- h) a polypeptide comprising an amino acid sequence of SEQ ID NO:34;
- i) a polypeptide comprising an amino acid sequence of SEQ ID NO:35;
- j) a polypeptide comprising an amino acid sequence of SEQ ID NO:36;
- k) a polypeptide comprising an amino acid sequence of SEQ ID NO:37;
- l) a polypeptide comprising an amino acid sequence of SEQ ID NO:38; and
- 15 m) a fragment of the polypeptide of (a), (b), (c), (d), (e), (f), (g), (h), (i), (j), (k), or (l)

wherein the expression of the mRNA encoding such polypeptide is altered in a T84 model of gut barrier function.

2. A purified polypeptide comprising an amino acid sequence that is at least 80% identical
20 to an amino acid sequence of claim 1.

3. An isolated DNA molecule encoding a purified polypeptide comprising an amino acid sequence that is at least 80% identical to an amino acid sequence of claim 1.

25 4. An isolated DNA molecule, wherein said isolated DNA molecule comprises a nucleotide sequence selected from the group consisting of:

- a) SEQ ID NO:1;
- b) SEQ ID NO:2;
- c) SEQ ID NO:3;

d) SEQ ID NO:4;
e) SEQ ID NO:5;
f) SEQ ID NO:6;
g) SEQ ID NO:7;
5 h) SEQ ID NO:8;
i) SEQ ID NO:9; and
j) SEQ ID NO:10 wherein the isolated DNA molecule has altered expression in a T84 model of gut barrier function.

10 5. An isolated DNA molecule, wherein said DNA molecule encodes a polypeptide comprising an amino acid sequence that is at least 80% identical to an amino acid sequence encoded by the DNA of claim 4.

15 6. An expression vector comprising the isolate DNA molecule according to claim 4.

7. A host cell transformed with the expression vector according to claim 5.

8. An isolated DNA molecule, wherein said isolated DNA molecule comprises a nucleotide sequence selected from the group consisting of: SEQ ID NO:1 - 26.

20 9. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a polynucleotide having a sequence selected from the group consisting of:

25 (a) a polynucleotide fragment of SEQ ID NO:1-26 or a polynucleotide which is hybridizable to SEQ ID NO:1-26;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 1-26 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO:1-26;

30 (c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 1-26 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO:1-26;

(e) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 1-26, having biological activity;

(f) a polynucleotide which is a variant of SEQ ID NO:1-26;

(g) a polynucleotide which is an allelic variant of SEQ ID NO:1-26;

5 (h) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 1-26;

(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A
10 residues or of only T residues.

10. The isolated nucleic acid molecule of claim 9, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.

15 11. The isolated nucleic acid molecule of claim 9, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a polypeptide chosen from the group consisting of:

(a) a polypeptide having the polypeptide sequence identified as a translation of SEQ ID NO: 1-26;

20 (b) a polypeptide having the polypeptide sequence of SEQ ID NO: 27-38; and

(c) a polypeptide encoded by the cDNA which is hybridizable to SEQ ID NO:1-26.

12. The isolated nucleic acid molecule of claim 9, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:1-26 or the cDNA sequence which is
25 hybridizable to SEQ ID NO:1-26.

13. The isolated nucleic acid molecule of claim 11 wherein the nucleotide sequence comprises sequential nucleotide deletions from sequence encoding either the C-terminus or the N-terminus.

30

14. The isolated nucleic acid molecule of claim 10 wherein the nucleotide sequence comprises sequential nucleotide deletions from sequence encoding either the C-terminus or the N-terminus.

15. A recombinant vector comprising the isolated nucleic acid molecule of claim 9.

16. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 9.

5

17. A recombinant host cell produced by the method of claim 16.

18. An isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence selected from the group consisting of:

10 (a) a polypeptide fragment of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(b) a polypeptide having the sequence of SEQ ID NO: 27-38;

(c) a polypeptide domain of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

15 (d) a polypeptide epitope of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(e) a secreted form of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(f) a full length protein of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

20 (g) a variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26;

(h) an allelic variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26; and

(i) a species homologue of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-26.

25

19. The isolated polypeptide of claim 18, wherein the full length polypeptide comprises sequential amino acid deletions from the C-terminus.

20. The isolated polypeptide of claim 18, wherein the mature polypeptide comprises sequential amino acid deletions from the C-terminus.

30

21. The isolated polypeptide of claim 18, wherein the full length polypeptide comprises sequential amino acid deletions from the N-terminus.

22. The isolated polypeptide of claim 18, wherein the mature polypeptide comprises sequential amino acid deletions from the N-terminus.

23. An isolated antibody that binds specifically to the isolated polypeptide of claim 18.

24. A recombinant host cell that expresses the isolated polypeptide of claim 18.

25. A method of making an isolated polypeptide comprising:

- (a) culturing the recombinant host cell of claim 24 under conditions such that said polypeptide is expressed; and
- (b) recovering said polypeptide.

26. The polypeptide produced by the method of claim 25.

27. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 18 or the polynucleotide of claim 9.

28. The method of claim 25 wherein the medical condition is irritable bowel disease.

29. A method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject comprising:

- (a) determining the presence or absence of a polynucleotide of claim 9; and
- (b) diagnosing irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or absence of said polynucleotide.

30. A method of diagnosing irritable bowel disease or a susceptibility to irritable bowel disease in a subject comprising:

- (a) determining the presence or amount of expression of the polypeptide of claim 18 in a biological sample; and
- (b) diagnosing irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or amount of expression of the polypeptide.

31. A method for identifying a binding partner to the polypeptide of claim 18 comprising:

- (a) contacting the polypeptide of claim 18 with a binding partner; and
- (b) determining whether the binding partner affects an activity of the polypeptide.

5

32. A method of identifying an activity in a biological assay, wherein the method comprises:

- (a) expressing the polynucleotide of SEQ ID NO:1-26 in a cell;
- (b) isolating the supernatant;
- (c) detecting an activity in a biological assay; and
- (d) identifying the polypeptide in the supernatant having the activity.

10

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Name: IMX4

CCGGTAAGTAAACAGTCAGAAAATTAGCATGAAAGCAGTTTAGCATTGGGAGGGAAGCACA
GATCTCTAGAGCTGTCCTGTCGCTGCCCAGGATTGACCTGTGTGTAAGTCCCAATAAACT
CACCTACTACCAAAAA

FIGURE 1

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Name: IMX 10

GAATTTAATACGACTCACTATAGGGAATTTGGCCCTCGAGGCCAAGAATTTCGGCACGAGG
CAACAACAACAACAAAAAAACTGAACATCTCCATATTACTGACACCCAATTCAAGAAA
CAAAATATTACAGCCCCCTTCCAGGATATTCCTGGGGTCTCTTCCATCTCTACTAACCCCT
GACTACAAACAGCCTCCACCTATTTTACCTGACATTGTACTTTATGAAAGCAGCAGTTCT
CAGATGGGGCTATTTTGCCCCCTGGGGACATTAGGGAGTATCTGGAGACACTGAGGGTTG
TGTCTACTTGGGGGGAGTTGTGTTACTGCATCCAGTGAGTCCAGGGATCCAGGGATGCCG
CTCAACATCCTGAAATGCACAGGGAACCCCCACACATAGAACAGAGAAATTGCTGAGCCA
AAATGTCAGCAGTGTCACAGCTGACACCCTGATATACACACTATCACACAGTATCTGCTC
TTTCGGGCTCAGGATCTTTTTTCAATTCTAATCATCTCATAGGAAACAGAAATGTCATTTAG
AGGTAGGTACAGTCCACAACAAAGAAGAACCTGAGTTTTTTTTTTTTTTTTTAATCAGCCT
GGTGCCTTTAGAGCTAGGATTTAGTTTCTATTCTTTCTGTCTCATTTTTCAAGTGATTTTT
TTCTTCAAATGGCATCTACTGGGCTCAAGAACTGGAGATCCCCACAAAGCTGAGATTCAC
ATGGGAATTTTGTACACACCCACACAGGTATACACTTCCATTTACATGCAGACATCCACC
CACAGATACACACATCCGGAGACCAAGACAGAACGCAAACCTGCCCCATAAAAGCACGGTT
CCCCAAACAGGAGAAACGCACCATTCCTCCAGGGAGGTATCTATTTGTTTAATTCAGCC
TCTGATAGTCAGGCTGTTGCCAAGCCCAGCTCTGAAACTCTTCCCCTCTAGGAAAGAAAG
ATGGATTTTTTCTTTACTCAAGAAATATAGATCTAAAAAAAAAAAAAAAAAAGTTGGCGG
CCGCAAGCTTATTCCTTTAGTGA

FIGURE 2

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Name: IMX 21

GGTACCGGGCCCCCCTCGAGGTCGACGGTATCGATAAGCTTGATATCGAATTTCGCGGGCC
GCTGAGAAATTAACTCCCCGGGGGCGCCGGGTTGACTGCGCTGCCTGGGGCCGGAGGTCTT
CTCCGGCCAGGGAGCGCTGTGGGAAGGGGCTCGAGCGGGCCAGGGCCAGGCGAGGCGGGG
GGGCGGGGGGGTTAGGGGACCGCGGGGGCTACTCTTGGGAGCGCCCCCTGTCCGGCTGGCTGC
GCGCCGGTTTTTAAATAGCATCTTTTCGGACTTGTCTTCGCGGCCCCAGTCCCCGACCTCGG
CGCTGCCTGGGCTCCTGCAGCCTCTCCCTAAGTCTTCTCCAAACGACCACCTCACGGATT
CCTTATGGATCGCAGCTCCAAGAGGAGGCAGGTGAAGCCTTTGGCAGCTTCTCTGCTGGA
AGCTCTTGATTATGATAGTTCAGATGACAGTGATTTTAAAGTTGGAGATGCCTCAGGACT
CGCTGATTCTTGAGAAGAGTCAAACTGGAGCTCTCAAAAAATGGACCATATTCTGATTT
GCTGTGTTTTGTCTGGGAGATAATAGTGAGGACGCTGATGAAATAATTCAGTGTGACAATT
GTGGCATTACAGTCCATGAAGGTTGTTATGGAGTTGATGGAGAGAGTGACTCTATTATGA
GTTTCAGCTTCTGAAAACCTCCACTGAACCTTGGTTTTGTGATGCCTGTAAATGTGGTGT
CTCCTAGCTGTGAACTGTGTCCTAATCAGGATGGAATTTTCAAGGAGACAGATGCTGGAA
GATGGGTTTCATATTGTTTTGTGCCCTGTATGTTCTTGAGTAGCCTTTGGAGATATTGACA
AATTACGACCAGTAACACTAACGGAAATGAACTATTCCAAATATGGTGCCAAGGAGTGTA
GCTTTTGTGAAGACCCTCGCTTTGCTAGAACTGGGGTTTTGCATTAGCTGTGATGCAGGGA
TGTGCAGAGCCTATTTCCATGTGACCTGTGCTCAAAAGGAAGGTCTGCTTTTCAGAGGCAG
CGGCGGAAGAGGATATAGCAGATCCATTCTTTGCTTATTGTAAAGCAACATGCAGATAGGT
TAGACAGAAAGTGGAAGAGAAAAAACTACTTGGCTCTACAGTCCTATTGTAAAATGTCTT
TGCAAGAGAGAGAGAAGCAACTATCACCAGAAGCACAGGCAAGGATCAATGCCCGGCTTC
AGCAGTATCGTGCCAAAGCAGAACTAGCTCGATCTACCAGACCCCAGGCCTGGGTTCCAA
GGGAAAAATTGCCCAGACCACTCACCAGCAGTGCTTCAGCTATTTCGTAAACTTATGCGGA
AAGCAGAACTCATGGGGATCAGTACAGATATCTTTCCAGTGGACAATTCAGATACTAGTT
CTAGTGTGGATGGAAGGAGAAAACATAAGCAACCAGCTCTCACTGCAGATTTTGTGAATT
ATTATTTTGTAGAGAAATATGCGCATGATTCAAATTCAGGAAAATATGGCTGAACAAAAGA
ATATAAAAGATAAATTAGAGAATGAACAAGAAAAGCTTCATGTAGAATATAATAAGCTAT
GTGAATCTTTAGAAAGAACTACAAAACCTGAATGGAAAACCTTCGAAGTGAAGGACAAAGGA
TATGGGCTTTACTAGGCAGAAATCACAGGGGCAGAAAGTTGAATATACCGGCAATTTTSCGAG
CACCCAAGGAGAGAAAACCAAGTAAAAAAGAAGGAGGCACACAAAAGACATCTACTCTTC
CTGCAGTACTTTATAGTTGTGGGATTTGTAAAGAAGAACCATGATCAGCATCTTCTTTTAT
TGTGTGATACCTGTAACTACATTACCATCTTGGATGTCTGGATCCTCCTCTTACAAGGA
TGCCAAGAAAGACCAAAAACAGTTATTGGCAGTGCTCGGAATGTGACCAGGCAGGGAGCA
GTGACATGGAAGCAGATATGGCCATGGAAACCCTACCAGATGGAACCAAAACGATCAAGGA
GGCAGATTAAGGAACCAGTGAAATTTGTTCCACAGGATGTGCCACCAGAACCCAAAGAGA
TTCCGATAAGAAACACGAGAACCAGAGGACGAAAACGAAGCTTCGTTCTTGAGGAAGAAA
AACATGAGGAAAGAGTTCTTAGAGAGAGAGAAGACAAAGACAGTCTGTGTTGCAAAAGAAGC
CCAAGGCTGAAGATTTAAGAACTGAATGTGCAACTTGCAAGGGAACTGGAGACAATGAAA
ATCTTGTGAGGTGTGATGAATGCAGACTCTGCTACCATTTTGGCTGTTTGGATCCTCCTT
TGAAAAAGTCTCCTAAACAGACAGGCTACGGATGGATATGTGAGGAATGTGATTCTTCAT
CTTCCAAGGAAGATGAAAATGAAGCTGAAAGAAAAAATATACTCAGGAGCTCAACATGG
AACAGAAAAATCCAAAGAAATAAAAGATTTTCTGTAGTGTTTTTGAAAAGTTTGCAGCTT
ATGTAATAGCAGATAAAATTTCTAATTGTAAAATGTTAAATTGAGCGGCCCGCGAATTCCT
GCAGCCCCGGGGGATCCACTAGTTCTAGAGCGGCCGCCACCGCGGTGGAGCTCCAGCT

FIGURE 3

4/23

Name: IMX 28

ATTAACCCTCACTAAAGGGGAACAAAAGCTGGAGCTCCACCGCGGTGGCGGGCCGCTCTAGA
ACTAGTGGATCCCCCGGGCTGCAGGAATTCGGCACGAGGTGCGCGGCTGCAACGGCAGCC
GCGGGAAGCTCGGGCCGGCAGGGTTTCCCCGCACGCTGGCGCCCCAGCTCCCCGGCGCGGAG
GCCGCTGTAAGTTTCGCTTTCCATTTCAGTGGAAAACGAAAGCTGGGCGGGGTGCCACGAG
CGCGGGGGCCAGACCAAGGCGGGGCCCGGAGCGGAACTTCGGTCCCAGCTCGGTCCCCGGCT
CAGTCCCGACGTGGAACTCAGCAGCGGAGGCTGGACGCTTGCATGGCGCTTGAGAGATTG
CATCGTGCCTGGCTCACATAAGCGCTTCCTGGAAGTGAAGTCGTGCTGTCCTGAACGCGG
GCCAGGCAGCTGCGGCCTGGGGGTTTTGGAGTGATCACGAATGAGCAAGGCGTTTGGGCT
CCTGAGGCAAATCTGTCAGTCCATCCTGGCTGAGTCCTCGCAGTCCCCGGCAGATCTTGA
AGAAAAGAAGGAAGAAGACAGCAACATGAAGAGAGAGCAGCCCAGAGAGCGTCCCAGGGC
CTGGGACTACCCTCATGGCCTGGTTGGTTTACACAACATTGGACAGACCTGCTGCCTTAA
CTCCTTGATTCAGGTGTTTCGTAATGAATGTGGACTTCACCAGGATATTGAAGAGGATCAC
GGTGCCCAGGGGAGCTGACGAGCAGAGGAGAAGCGTCCCTTTCCAGATGCTTCTGCTGCT
GGAGAAGATGCAGGACAGCCGGCAGAAAGCAGTGCGGGCCCCCTGGAGCTGGCCTACTGCCT
GCAGAAGTGCAACGTGCCCTTGTTTGTCCAACATGATGCTGCCCAACTGTACCTCAAAC
CTGGAACCTGATTAAGGACCAGATCACTGATGTGCACTTGGTGGAGAGACTGCAGGCCCT
GTATATGATCCGGGTGAAGGACTCCTTGATTTGCGTTGACTGTGCCATGGAGAGTAGCAG
AAACAGCAGCATGCTCACCCTCCCACCTTTCTCTTTTTTGATGTGGACTCAAAGCCCCCTGAA
GACACTGGAGGACGCCCTGCACTGCTTCTTCCAGCCCAGGGAGTTATCAAGCAAAGCAA
GTGCTTCTGTGAGAACTGTGGGAAGAAGACCCGTGGGAAACAGGTCTTGAAGCTGACCCA
TTTGCCCCAGACCCTGACAATCCACCTCATGCGATTCTCCATCAGGAATTACACAGACGAG
AAAGATCTGCCACTCCCTGTACTTCCCCCAGAGCTTGGATTTTCAGCCAGATCCTTCCAAT
GAAGCGAGAGTCTTGTGATGCTGAGGAGCAGTCTGGAGGGCAGTATGAGCTTTTTTGCTGT
GATTGCGCACGTGGGAATGGCAGACTCCGGTCATTACTGTGTCTACATCCGGAATGCTGT
GGATGGAAAATGGTTCTGCTTCAATGACTCCAATATTTGCTTGGTGTCTTGGGAAGACAT
CCAGTGTACCTACGGAAATCCTAACTACCACTGGCAGGAAACTGCATATCTTCTGGTTTA
CATGAAGATGGAGTGCTAATGGAAATGCCCAAACCTTCAGAGATTGACACGCTGTCATT
TTCCATTTCCGTTTCTGGATCTACGGAGTCTTCTAAGAGATTTTGCAATGAGGAGAAGCA
TTGTTTTCAAACCTATATAACTGAGCCTTATTTATAATTAGGGATATTATCAAAATATGTA
ACCATGAGGCCCCCTCAGGTCCTGATCAGTCAGAATGGATGCTTTCCACAGCAGACCCGGC
CATGTGGCTGCTCGGTCTTGGGTGCTCGCTGCTGTGCAAGACATTAGCCCTTTAGTTATG
AGCCTGTGGGAACCTTCAGGGGTTCACAGTGGGGAGAGCAGTGGCAGTGGGAGGCATCTGG
GGGCCAAAGGTCAGTGGCAGGGGGTATTTTCAGTATTATACAACCTGCTGTGACACAGACTTG
TATACTGGCTGAATATCAGTGCTGTTTGTAAATTTTTCACCTTTGAGAACCAACATTAATTC
CATATGAAAAAAAAAAAAAAAAAAAAA

FIGURE 4

5/23

Name: IMX 32

GCGGCCGCTCTAGAACTAGTGGATCCCCCGGGCTGCAGGAATTCGCGGCCGCTAAATGAA
CTCCCATAGAGTCTAGACACCATAGAACTCATACCAGGAATCACAAAGTCTCTAAATTT
CCAAAGTTAACTGGAAATATTACAAACTGCAGAATAATTCCAGGCCAAAATATGTTAAAT
TCATAACATGATGTATATCAAAGGAAAAAAGGACATGTGGAAATGACACATTATCTTCAG
TGTATAAAATATTCATTTATGTGAAGTTTCTTGGAAGGCTACACTACTATTACTGGTTT
CCGTCTGATGTTTGAGATCTGTTGATTTTATGCTTTTCTTACAGGCCTTTCATTATGATC
TTTGGGAAGGAATCAATAAAATGATAGGGCCTACTTCATTAGGTGTGGTTCATTCCTATT
CATGCTCCCTGGAAGAACAAGAATGCTGAATTTTGAAATTTAATATTGTATGAATTAGCA
TCAGGGAGAGGTGGAGAAAAATACAAAACCTAAAAGTCATGCTTATTGTGTTCAGTGTGCC
CTTCTCCAGAGGGGCCACTGGCTTATAGGAAAGGATTGCTGCTCTACCAGTTGACCAGGAG
ATGGCACGCCAGGACATTAAGACACTGGAGTTTTGTTTCGTTTTTTTTTTTTTTTTTTGAG
ATGGAGTCTCGCTCTCTTGACAGGCAGGAGTACAGTGGTGCGATCTCGGCTCACTGCAAA
CTCCGCCTCCCGGGTTCAAGTGATTCTCCTGCCTCGGCCTCCCGAGTAGCTGGGACTACA
GGCGTGTGCCACCACCCCCAGCTAACTTTTGTATTTTATAGTAGAGACAGGGTTTCACCAT
GTTGGCCAGGATGGTCTCAATCTCTTGACCTCATGATCCGCCCGCCTCCGCCTTCCAAAG
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TTTTTAAAAGCAATAGCTGTTGTATATCAATGATAAATGAAAAATTAGTTATTCTTGTA
ATTGAAGAAAGAATGGTTATCATAGAGGGTAGTTCAAGTAAAAGAACCAGGGCTGGGTGT
GGTGGCTCACGTTCTGTAATCCCTGTACTTTGGGAGGCCAAGGCAGATGGATCTCTTGAG
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AAAATCACTTGAACCCGGGGGACGGAGGTTGCAGTGAGCTGAGATCGCACCACTGCTCGC
CAGCCTGGGCAACAGAGTGAGACTCTGCCTCAAAAAAAAACCAAACCAAACCAAAGAACC
AGAATAGCATGTGCACATATACACAGACGTTTCACAACTGGCATTATGTTTTGCTACTGT
TTTATTTACAATGTATCACAAGTTTTATGCTTTAATAAAATTTAATCATAACTTCAAAAA
AAAAAAAAAAAAAAAAAGCGGCCGCGAATT

FIGURE 5

6/23

Name: IMX 39

GAATTCGGCACGAGGAAAACATTTGCCCCCTTGCAGAAGATCACCCCTTAGTTCTTCCTCGG
AAGAGTATCAGAAGGTCTGGAACCTCTTTAACCGCACGCTGCCTTTCTACTTTGTTTCAGA
AGATTGAGCGAGTACAGAACCTGGCCCTCTGGGAAGTCTACCAGTGGTGCGTTGGGGCTC
GCTCTTGGTGGGCTGGTGACTCTGTCCCTTCACACCACTGGCTGGTTGCCACATGTGGCC
CGGGTTTCCAGGAAAAGCAGAGCGGCAGTTAGGGCTGCCATGTGCTGGGAGCTGTGTGTC
TGCTCTCCTTCGTCCGCTCCCCCAGGGCAGTGTGGTAGCACATCCCATTGTAGAGATGAG
GGCACCGAGGCTTCCTGGAGCATACCACCTGGTCCCGTTTCATGAGTGGTGGCAAAGCTAG
CACTCTCACTTGTCATTCTGCCTTCCTGGAGACCAGTGGGATGGGTGAGTACAGCCAC
CACACCATTAGCCCCAGGAACATAAGGCTGTGGCTAGACAGCAGGGGTCTCAGGTTTATA
CATGAGGACTGGCTTGTCCTTGAGCACCCACTCACCTGTCTATGTGGGGAGGAATCCTAC
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GTAATGTGTACTCTGTGCACAGTGATGAAGTCTGGGAATGGGAGAGGGGAGAAGGATGGG
CACCCACTGACCAGCAGCCTGAAAATTCCTACAGCATCCCAGGGCTCAGCTCCATGCAGG
AGCAAGGTGGGGGTGGGGTTGGGGGAAATGTTACCCATTTTCCAAGGGCTGCTCTGCTTT
TGGAGTCCAGGGAACCGCTGCTGTCTGGAGCTGTGGAGGGAGGGTTTTTCACCCAGCTCCC
ACGATCCCCCTTCTTTTCCACACCCTGGCTTGTTGGCTGGAGCCTTACAGGCCTAGTCAGG
GTAGCCTGTGACCTGCGTCTCTTGGTCCCAGGACACTTTTGGAATTTTGGAAAAATGTGT
TGTTTTGCATCAGGCCGGCTGTATTTGGTGGCCGGCACACTCTGCCCCCAGCACACATTC
TTCTGTGATTCTAGGCAAAAAGGACAGATGCAGAAGCAGAACGGAGGGAAGGCCGTGGAC
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TTTGACTGGCGGGTCTGTGGTGTTCATGGCACTTCCTACGGCAAGGGGAGCTACTTTGCC
CGAGATGCTGCATATTCCCACCACTACAGCAAATCCGACACGCAGACCCACACGATGTTT
CTGGCCCCGGGTGCTGGTGGGCGAGTTCGTGAGGGGCAATGCCTCCTTTGTCCGTCCGCCG
GCCAAGGAGGGCTGGAGCAACGCCTTCTATGATAGCTGCGTGAACAGTGTGTCCGACCCC
TCCATCTTTGTGATCTTTGAGAAACACCAGGTCTACCCAGAGTATGTCATCCAGTACACC
ACCTCCTCCAAGCCCTCGGTACACCCCTCCATCCTGCTGGCCTTGGGCTCCCTGTTTCAGC
AGCCGACAGTGAGCGCACAGGAGTGTTCCAGGCCTTTCACCTGCTCTGCCTTGAAATGGC
TATTTGGGCCCTTTCCTTTTCTTTTAAACAGAACTTTTAATGAACTGTTCTCTTAACAT
TGACCTCTCAATGAAGTTATGTTCTTAATCTCTTGCTAATAATGATTTTTTACTTTTAAAGT
CACTTTTGGGTTCACTAGTGGATTAACCAGAAGTGATTGTAGTTGAGTCCAGTTTTGCTT
TTTAATAATGTGTTGAAGTTTTAGTTTTTACTCTTTGTTGACTTTGCTGCTTATTGGCAC
CAGGGACAGAGTTTCTAGATACAATTTTATGGATTGGTTTTTAATTTTATGAGTTTGTCT
CTGCAGTGATTGGTTTTCTCAGAGTCTCATGGCATCATAGTTTTTCCAGAATGACACAGT
AGCCACCGGTGGATGACAGCCACGGGCGGCACAGTCACTTCTGCCTGTTGCTCTGACAC
CAACCCAGGCAGCTCTGCTGTGGCTTCTCCTGGGCTCTGGCATTAGTTGGTCTGTGTGAC
ATTGTCAGAACAGGTGGCTGCTGTGTGGTGCCATCGAGTCCCTGCTGGTTCCCTTGTCC
TGGGAGGGTCACCCATTGCCCAAGGAAGTGCATCCACCTGGCAGGTGACCTGGAGGAGTA
GCTTCCCCGAGGACCCCCAGGCTTGGCCTGTGATTGCGCAAACCCACATTTCTTAAGCAC
ACTGGACACCCTTCGAGTGTGGGTTTTTAACATCCCTGTGAGATTGAATACTTGTGCCACA
CATGTCACAAAAGAGTATGGAAATAAAAGAAAATTTATCCGAAAAAAAAAAAAAAAAAAT
GAGCGGCCCGC

FIGURE 6

7/23

Name: IMX 40

CCGGTCTATGGCATTAAACCCTCACTTAACTTTTCAGCCTGCCAGCCTGCCCTATGGATT
CGGACTTGCCAGCCACACAATTCCTTAAAATAAATCTCTCCGTCTCATAAAAA

FIGURE 7

8/23

Name: IMX 42

CCGGGAGCTGTGAAGGGAACGTGAGGGGGCGGCGTAGTGGAGACCCACGGCAGGCCTGAA
GAAGAGCGGCGGCCGAGCCCGCCTTCCCTGCACCATGCTCATAGAGGATGTGGATGCCCT
CAAGTCCTGGCTGGCCAAGTTACTGGAGCCGATATGTGATGCTGATCCTTCAGCCTTAGC
CAACTATGTTGTAGCACTGGTCAAGAAGGACAAACCTGAGAAAGAATTAAAAGCCTTTTG
TGCTGATCAACTTGATGTCTTTTTTACAAAAGAACTTCAGGTTTTGTGGACAAACTATT
TGAAAGTCTCTATACTAAGAACTACCTTCCACTTTTGGAACCAGTAAAGCCTGAGCCAAA
ACCACTAGCCCAAGAAAAA

FIGURE 8

9/23

Name: IMX 44

ATGGATAGTCGCCACACCTTTGCCCCCTGCTGCGATGACCCTGTGCGCCACTTCTGCTGTTTC
CTGCCACCGCTGCTGCTGCTGCTGGACGTCCCCACGGCGGGCGGTGCAGGCGTCCCCCTCTG
CAAGCGTTAGACTTCTTTGGGAATGGGCCACCAGTTAACTACAAGACAGGCAATCTATAC
CTGCGGGGGCCCCCTGAAGAAGTCCAATGCACCGCTTGTCAATGTGACCCTCTACTATGAA
GCACTGTGCGGTGGCTGCCGAGCCTTCCTGATCCGGGAGCTCTTCCCAACATGGCTGTTG
GTCATGGAGATCCTCAATGTCACGCTGGTGCCCTACGGAAACGCACAGGAACAAAATGTC
AGTGGCAGGTGGGAGTTCAAGTGCCAGCATGGAGAAGAGGAGTGCAAATTCAACAAGGTG
GAGGCCTGCGTGTTGGATGAACTTGACATGGAGCTAGCCTTCCTGACCATTGTCTGCATG
GAAGAGTTTGAGGACATGGAGAGAAGTCTGCCACTATGCCTGCAGCTCTACGCCCCAGGG
CTGTGCGCCAGACACTATCATGGAGTGTGCAATGGGGGACCCCGGCATGCAGCTCATGCAC
GCCAACGCCCAGCGGACAGATGCTCTCCAGCCACCACACGAGTATGTGCCCTGGGTCACC
GTCAATGGGAAACCCTTGGAAGATCAGACCCAGCTCCTTACCCTTGTCTGCCAGTTGTAC
CAGGGCAAGAAGCCGGATGTCTGCCCTTCCTCAACCAGCTCCCTCAGGAGTGTTTGCTTC
AAGTGATGGCCGGTGAGCTGCGGAGAGCTCATGGAAGGCGAGTGGGAACCCGGCTGCCTG
CCTTTTTTTTTCTGATCCAGACCCTCGGCACCTGCTACTTACCAACTGGAAAATTTTATGC
ATCCCATGAAGCCCAGATACACAAAATTCCACCCCATGATCAAGAATCCTGCTCCACTAA
GAATGGTGCTAAAGTAAACTAGTTTAATAAGCAAAAAAAAAAAAAAAAAAATTCCTGCG
GCCGC

FIGURE 9

10/23

Name: IMX 56

CCGGGATATCGCCACTGCACTCCAGCCTGGGTGACGGAGCGAGACTCCGTCTCAGAAAAA

FIGURE 10

11/23

Name: IMX4

MHWEEAQISRVLSPRIDLCVSPNKLTYSRK

FIGURE 11

12/23

Name: IMX 10

MEFNTTHYREFGPRGQEFGRQQQQQKKTEHLHITDTQFKKQONITAPSRIFLGSLPSLLT
PDYKQPPPISPDIVLYESSSSQMGLFCPLGTLGSIWRH*

FIGURE 12

13/23

Name: IMX 21

MI VQMTVILKLEMPQDSLILEKSQNWSSQKMDHILICCVCLGDNSEDADEIIQCDNCGIT
VHEGCGVDGESDSIMSSASENSTEPWFCDACKCGVSPSCELCPNQDGI FKETDAGRWH
IVCALYVPGVAFGDIDKLRPVTLTENMYSKYGAKECSFCEDPRFARTGVCISCDAGMCRA
YFHVTCAQKEGLLSEAAAEEDIADPFFAYCKQHADRDRKWKRRKNYLALQSYCKMSLQER
EKQLSPEAQARINARLOQYRAKAELARSTRPQAWVPREKLPRPLTSSASAIRKLMRKAEL
MGISTDIFPVDNSDTSSSV DGRRKHKQPALTADFNYYFERNMRMIQIQENMAEQKNIKD
KLENEQEKLHVEYNKLCESLEELQNLNGKLRSEGQGIWALLGRITGQKLNIPAILRAPKE
RKPSKKEGGTQKTSTLPAVLYSCGICKKNHDQHLLLLCDTCKLHYHLGCLDPPLTRMPRK
TKNSYWQCSECDQAGSSDMEADMAMETLPDGTKRSRROIKEPVKFVPQDVPPEPKKIPR
NTRTRGRKRSFVPEEEKHEERVPRERRQRQSVLQKKPKAEDLRTECATCKGTGDNENLVR
CDECRLCYHFGCLDPPLKKSPKQTGYGWICQECDSSSSKEDENEAERKNISQELNMEQKN
PKK

FIGURE 13

14/23

Name: IMX 28

MSKAFGLLRQICQSILAESSQSPADLEEKKEEDSNMKREQPRERPRAWDYPHGLVGLHNI
GQTCCLNSLIQVFVMNVDFTRILKRITVPRGADEQRRSVPFQMLLLEKMQDSRQKAVRP
LELAYCLQKCNVPLFVQHDAQAQLYLKLWNLIKDQITDVHLVERLQALYMIRVKDSLICVD
CAMESSRNSSMLTLPLSLFDVDSKPLKTLEDALHCFFQPRELSSKSKCFCENCCKTRGK
QVLKLTHLPQTLTIHLMRFSIRNSQTRKICHSLYFPQSLDFSQILPMKRESCDAEEQSGG
QYELFAVIAHVGMADSGHYCVYIRNAVDGKWFCFNDSNICLVSWEDIQCTYGNPNYHWQE
TAYLLVYMKMEC*

FIGURE 14

15/23

Name: IMX 32

MAAALLPSGQNWHTGFILESNLTNVMKVVRLEFIKCPCLWGHEKIHTESIKNVLNMERPL
SNSDVMKVVF*

FIGURE 15

16/23

Name: IMX 39

MCTLCTVMKSGNGRGEKDGHP LTSSLKIPTASQGSAPCRSKVGVGLGEMLP I FQGLLCFW
SPGNRCCLELWREGFHPAPT I PLLFHTLACGWSLTGLVRVACDLRLLVPGHFWNFGKMCC
FASGRLYLVAGTLC PQHTFFCDSRQKGQMQKQNGGKAVDERQLFHGTSAI FVDAICQQNF
DWRVCGVHGTSYGKGSYFARDAAYSHHYSKSDTQTHTMFLARVLVGEFVRGNASFVRPPA
KEGWSNAFYDSCVNSVSDPSIFVIFEKHQVYPEYVIQYTTSSKPSVTPSILLALGSLFSS
RQ

FIGURE 16

17/23

Name: IMX 40

MPVYGINPHLTFQPASLPYGFRTCQPHNSLK*

FIGURE 17

18/23

Name: IMX 42

MLIEDVDALKSWLAKLLEPICDADPSALANYVVALVKKDKPEKELKAFCADQLE:FLQKE
TSGFVDKLFESLYTKNYLPLLEPVKPEPKPLAQEK*

FIGURE 18

19/23

Name: IMX 44

(MD) SRHTFAPAAMTLSPLLLFLPPLLLLLLDVPTAAVQASPLQALDFFGNGPPVNYKTGN
LYLRGPLKKSNAPLVNVTLYYEALCGGCRAFLIRELFPTWLLVMEILNVTLPYGNAAEQ
NVSGRWEFKCQHGEEECKFNKVEACVLDELDMELAFLTIVCMEEFEDMERSLPLCLQLYA
PGLSPDTIMECAMGDPGMQLMHANAQRTDALQPPHEYVPWVTVNGKPLEDQTQLLTLCQ
LYQGKKPDVCPSSSTSSLRVCFK

FIGURE 19

20/23

Name: IMX 56
MPGYRHCTPAWVTERDSVSEK*

FIGURE 20

Match to cDNA gb|AF019225|AF019225 Homo sapiens apolipoprotein L mRNA, complete cds
 Length = 1279
 Plus Strand HSPs:

Score = 2078 (311.8 bits), Expect = 6.0e-88, P = 6.0e-88
 Identities = 432/451 (95%), Positives = 432/451 (95%), Strand = Plus / Plus

Query: 168 ATGAGTGCACCTTTTCCTTGNNTNANTNANONCAGAGGAGAGCTGGAGCGAGGGGTGCAA 227
 Sbjct: 1 ATGAGTGCACCTTTTCCTTGGTGTGAGAGCTGGAGCGAGGGGTGCAA 60

Query: 228 CAAAACGTTCCAAAGTGGACAGATACCTGGAGATCCTCAAGTAAGCCCTCGGTGACTGG 287
 Sbjct: 61 CAAAACGTTCCAAAGTGGACAGATACCTGGAGATCCTCAAGTAAGCCCTCGGTGACTGG 120

Query: 288 GCTGCTGGCACCATTGGACCCAGAGAGCAGTATCTTTATTGAGGATGCCATTAAAGTATTC 347
 Sbjct: 121 GCTGCTGGCACCATTGGACCCAGAGAGCAGTATCTTTATTGAGGATGCCATTAAAGTATTC 180

Query: 348 AAGGAAAGTGGAGCACACAGAACTGCTACTCTGCTGACTGATTAATGAGGCCCTGGAAAC 407
 Sbjct: 181 AAGGAAAGTGGAGCACACAGAACTGCTACTCTGCTGACTGATTAATGAGGCCCTGGAAAC 240

Query: 408 GGATTCGTGGCTGCTGTAAGTCCCGAGGATGAGGCGATGAGCTCCGTAAAGCTCTG 467
 Sbjct: 241 GGATTCGTGGCTGCTGTAAGTCCCGAGGATGAGGCGATGAGCTCCGTAAAGCTCTG 300

Query: 468 GACAACTTGCAGACAAATGATCATGAAAGACAAAACTGGCACGATAAAGGCCAGCAG 527
 Sbjct: 301 GACAACTTGCAGACAAATGATCATGAAAGACAAAACTGGCACGATAAAGGCCAGCAG 360

Query: 528 TACAGAACTGGTTTCTGAAAGAGTTTCTCGGGTGAAAGTAAGCTTGAGGATAACA 587
 Sbjct: 361 TACAGAACTGGTTTCTGAAAAGTTTCTCGGGTGAAAGTAAGCTTGAGGATAACA 418

Query: 588 TAAAGAAAGCTTCGGTGCCCTTGCAAAATGGG 618
 Sbjct: 419 CAAAGAGGCT-CCGTGCCCTTGCG-CAATGGG 447

FIGURE 21

22/23

Match to exons on PAC clone that carries ApoL gene:

>emb|282215|HS6802 Homo sapiens DNA sequence from PAC 6802 on chromosome 22.

Contains apolipoprotein L, myosin heavy chain, ESTs, CA repeat, STS

and GSS, complete sequence [Homo sapiens]

Length = 139,389

Match of Query: 2

ACGAGCTGTCTGGTTATTATACAGACGCATAACTGGAGGTGGGATCCACACAGCTCAGAA 61

|||||
|||||

Sbjct: 21414

AGGACCTGTCTGGTTATTATACAGACGCATAACTGGAGGTGGGATCCACACAGCTCAGAA 21473

Query: 62 CAGCTGGATCTTGCTCAGTCTCTGCCAGGGGAAGATTCCTTGG 104

|||||
Sbjct: 21474 CAGCTGGATCTTGCTCAGTCTCTGCCAGGGGAAGATTCCTTGG 21516

Query: 103

GGAGGAGGCCCTGCAGCGACATGGAGGGAGCTGCTTTGCTGAGAGTCTCTGTCCTCTGCA 162

GGAGGAGGCCCTGCAGCGACATGGAGGGAGCTGCTTTGCTGAGAGTCTCTGTCCTCTGCA

Sbjct: 23232

GGAGGAGGCCCTGCAGCGACATGGAGGGAGCTGCTTTGCTGAGAGTCTCTGTCCTCTGCA 23291

Query: 163 TCTGGATGAG 172

TCTGG TGAG

Sbjct: 23292 TCTGG-TGAG 23300

Query: 121 ACATGGAGGGAGCTGCTTTGCTGAGAGTCTCTGTCCT-
CTGCATCTGGATGAGTGCACTT 179

A ATGG G T CT TG T A TCTC TC T C CA C

GGATGAGTGCACTT

Sbjct: 25333 AGATGGCTGCCCGTCTCTGATTATCTTCTCC-

TCATACCCCAACAGGATGAGTGCACTT 25391

Query: 180 TTCCTTGNTNTNGNANTNANGHCAGAGGAAGCTGGAGCGAGG 221

TTCCTTG T T G A T A G CAGAGGAAGCTGGAGCGAGG

Sbjct: 25392 TTCCTTGGTGTGGGAGTGAGGGCAGAGGAAGCTGGAGCGAGG 25433

Query: 217

CGAGGGTGCAACAAAACGTTCCAAGTGGGACAGATACTGGAGATCCTCAAAGTAAGCCCC 276

|||||
|||||

Sbjct: 25611

CAAGGGTGCAACAAAACGTTCCAAGTGGGACAGATACTGGAGATCCTCAAAGTAAGCCCC 25670

Query: 277 TCGGTGACTGGGCTGCTGGCACCATGGACCCAGAGAG 313

|||||
Sbjct: 25671 TCGGTGACTGGGCTGCTGGCACCATGGACCCAGGTAG 25707

FIGURE 22

23/23

Query: 304
ACCCAGAGAGCAGTATCTTTATTGAGGATGCCATTAAGTATTTCAAGGAAAAAGTGAGCA 363
|||||
Sbjct: 29886
AACTAGAGAGCAGTATCTTTATTGAGGATGCCATTAAGTATTTCAAGGAAAAAGTGAGCA 29945

Query: 364
CACAGAATCTGCTACTCCTGCTGACTGATAATGAGGCCTGGAACGGATTCGTGGCTGCTG 423
|||||
Sbjct: 29946
CACAGAATCTGCTACTCCTGCTGACTGATAATGAGGCCTGGAACGGATTCGTGGCTGCTG 30005

Query: 424 CTGAACTGCCCAGGAATG 441
|||||
Sbjct: 30006 CTGAACTGCCCAGGTAAG 30023

Query: 430
TGCCCAGGAATGAGGCAGATGAGCTCCGTAAAGCTCTGGACAACCTTGCAAGACAAATGA 489
TG
CAGGAATGAGGCAGATGAGCTCCGTAAAGCTCTGGACAACCTTGCAAGACAAATGA
Sbjct: 33440
TGTGCAGGAATGAGGCAGATGAGCTCCGTAAAGCTCTGGACAACCTTGCAAGACAAATGA 33499

Query: 490
TCATGAAAGACAAAAACTGGCACGATAAAGGCCAGCAGTACAGAACTGGTTTCTGAAA 549
TCATGAAAGACAAAAACTGGCACGATAAAGGCCAGCAGTACAGAACTGGTTTCTGAAA
Sbjct: 33500
TCATGAAAGACAAAAACTGGCACGATAAAGGCCAGCAGTACAGAACTGGTTTCTGAAA- 33558

Query: 550
GAGTTTCCTCGGGTGAAAAAGTAAGCTTGAGGATAACATAAGAAAGCTTCCGTGCCCTTG 609
GAGTTTCCTCGG TGAAAA GT AGCTTGAGGATAACATAAGAA GCT
CCGTGCCCTTG
Sbjct: 33559 GAGTTTCCTCGGTTGAAAA-GTGAGCTTGAGGATAACATAAGAAGGCT-
CCGTGCCCTTG 33616

Query: 610 CAANATGGG 618
CA ATGGG
Sbjct: 33617 CA-GATGGG 33624

FIGURE 22, CONTINUED

SEQUENCE LISTING

<110> Baum, Peter
 DuBose, Robert
 5 Sims, John E
 Youakim, Adel
 Hasel, Karl W
 Hilbush, Brian S

10 <120> Novel DNAs and Polypeptides
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 cacctactca ccaaaaaa 137

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 actagttcta gtgtggatgg aaggagaaaa cataagcaac cagctctcac tgcagatttt 240
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 caaaaa 306

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 caaaaa 366

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<213> Homo sapiens

15 <400> 5

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 gataaatgaa aaattagtta ttcttgtaaa ttgaagaaag aatgggttatc atagagggta 180
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Glu Asn Ser Thr Glu Pro Trp Phe Cys Asp Ala Cys Lys Cys Gly Val
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NOVEL DNAs AND POLYPEPTIDES**Reference to Related Application**

This application claims the benefit of U.S. Provisional Application S.N. 60/107,821, filed November 10, 1998, which is incorporated herein by reference.

BACKGROUND OF THE INVENTION**Field of the Invention**

The invention is directed to purified and isolated novel polypeptide molecules and fragments thereof, the nucleic acid molecules encoding such polypeptides, processes for production of recombinant forms of such polypeptide molecules, antibodies generated against such polypeptide molecules, fragmented peptides derived from these polypeptide molecules, and uses thereof. In particular, the invention is directed to the use of the nucleic acid and polypeptide molecules in the study of inflammatory bowel diseases (IBD)

Description of Related Art

Damage to the intestinal epithelial barrier is a hallmark of inflammatory bowel diseases (IBD). Examples of inflammatory bowel diseases include ileitis, Crohn's disease (CD), which can affect the whole digestive tract from mouth to anus, and ulcerative colitis (UC), which affects only the large intestine. Studying the factors that influence the integrity of the epithelial barrier *in vivo* is a difficult task for a number of reasons that include the complexity of the tissue itself (there are numerous cell types in the gut including epithelial, stromal, endocrine, neuronal and hematopoietic) and the technical problems associated with tissue manipulation in animals or in isolated organs. As a result of these issues, a number of *in vitro* models of epithelial barrier function have been developed over the years. The best characterized of these models is the T84 intestinal epithelial barrier system (Dharmasathaphorn et al., *Am. J. Physiol.*, 246:G204-G208, 1984 and Madara et al., *J. Cell Biol.*, 101:2124-2133, 1985). T84 cells, while derived from a human colonic adenocarcinoma, have retained many of the properties associated with normal colonic crypt cells. T84 cells form polarized monolayers that exhibit high electrical resistance and vectorial fluid and chloride secretion reminiscent of colonic crypts *in vivo*. These properties

are directly dependent on a complex of proteins referred to as tight junctions and T84 cells express many of the known members of this complex (Youakim et al., *Submitted for publication*, 1998). These cells also respond to proinflammatory cytokines, such as interferon-gamma (INF-gamma), by decreasing barrier function (Youakim et al., *Submitted for publication*, 1998; Madara et al., *J. Clin. Invest.*, 83:724-727, 1989; and Adams et al., *J. Immunol.*, 150:2356-2363, 1993) and by up-regulating MHC Class II molecules and antigen presenting activity (Hershberg et al., *J. Clin. Invest.*, 102:792-803, 1998). This model system is used to examine how the epithelial barrier is regulated by various agents such as interferon-gamma and cells of the immune system (e.g., neutrophils). The T84 model assists in the elucidation of the mechanism of barrier breakdown and recovery in response to these agents and in the identification of proteins (and genes) that may prevent barrier breakdown or stimulate barrier recovery.

Given the important function of the epithelial gut barrier and despite the growing body of knowledge, there is a need in the art for the discovery, the identification, and the elucidation of the roles of new proteins involved in gut barrier function and IBD.

The identification of the primary structure, or sequence, of an unknown protein is the culmination of an arduous process of experimentation. In order to identify an unknown protein, the investigator can rely upon a comparison of the unknown protein to known peptides using a variety of techniques known to those skilled in the art. For instance, proteins are routinely analyzed using techniques such as electrophoresis, sedimentation, chromatography, sequencing and mass spectrometry.

In particular, comparison of an unknown protein to polypeptides of known molecular weight allows a determination of the apparent molecular weight of the unknown protein (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)). Protein molecular weight standards are commercially available to assist in the estimation of molecular weights of unknown protein (New England Biolabs Inc. Catalog:130-131, 1995; J. L. Hartley, U.S. Patent No. 5,449,758). However, the molecular weight standards may not correspond closely enough in size to the unknown protein to allow an accurate estimation of apparent molecular weight. The difficulty in estimation of molecular weight is compounded in the case of proteins that are subjected to fragmentation by chemical or enzymatic means, modified by post-translational modification or processing, and/or associated with other proteins in non-covalent complexes.

In addition, the unique nature of the composition of a protein with regard to its specific amino acid constituents results in unique positioning of cleavage sites within the protein. Specific fragmentation of a protein by chemical or enzymatic cleavage results in a unique "peptide fingerprint" (D. W. Cleveland et al., *J. Biol. Chem.* 252:1102-1106, 1977; M. Brown et al., *J. Gen. Virol.* 50:309-316, 1980). Consequently, cleavage at specific sites results in reproducible fragmentation of a given protein into peptides of precise molecular weights. Furthermore, these peptides possess unique charge characteristics that determine the isoelectric pH of the peptide. These unique characteristics can be exploited using a variety of electrophoretic and other techniques (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)).

Fragmentation of proteins is further employed for amino acid composition analysis and protein sequencing (P. Matsudaira, *J. Biol. Chem.* 262:10035-10038, 1987; C. Eckerskorn et al., *Electrophoresis* 1988, 9:830-838, 1988), particularly the production of fragments from proteins with a "blocked" N-terminus. In addition, fragmented proteins can be used for immunization, for affinity selection (R. A. Brown, U.S. Patent No. 5,151,412), for determination of modification sites (e.g. phosphorylation), for generation of active biological compounds (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 300-301 (Prentice Hall, 6d ed. 1991)), and for differentiation of homologous proteins (M. Brown et al., *J. Gen. Virol.* 50:309-316, 1980).

In addition, when a peptide fingerprint of an unknown protein is obtained, it can be compared to a database of known proteins to assist in the identification of the unknown protein using mass spectrometry (W.J. Henzel et al., *Proc. Natl. Acad. Sci. USA* 90:5011-5015, 1993; D. Fenyo et al., *Electrophoresis* 19:998-1005, 1998). A variety of computer software programs to facilitate these comparisons are accessible via the Internet, such as Protein Prospector (Internet site: prospector.uscf.edu), MultiIdent (Internet site: www.expasy.ch/sprot/multiident.html), PeptideSearch (Internet site: www.mann.embl-heidelberg.de...deSearch/FR_PeptideSearch_Form.html), and ProFound (Internet site: www.chait-sgi.rockefeller.edu/cgi-bin/prot-id-frag.html). These programs allow the user to specify the cleavage agent and the molecular weights of the fragmented peptides within a designated tolerance. The programs compare these molecular weights to protein molecular weight information stored in databases to assist in determining the identity of the unknown protein. Accurate information concerning the number of fragmented peptides and the precise molecular weight of those peptides is required for accurate identification. Therefore,

increasing the accuracy in determining the number of fragmented peptides and their molecular weight should result in enhanced likelihood of success in the identification of unknown proteins.

In addition, peptide digests of unknown proteins can be sequenced using tandem mass spectrometry (MS/MS), and the resulting sequence searched against databases (J.K. Eng, et al., *J. Am. Soc. Mass Spec.* 5:976-989 (1994); M. Mann and M. Wilm, *Anal. Chem.* 66:4390-4399 (1994); J.A. Taylor and R.S. Johnson, *Rapid Comm. Mass Spec.* 11:1067-1075 (1997)). Searching programs that can be used in this process exist on the Internet, such as Lutefisk 97 (Internet site: www.lsbc.com:70/Lutefisk97.html), and the Protein Prospector, Peptide Search and ProFound programs described above. Therefore, adding the sequence of a gene and its predicted protein sequence and peptide fragments to a sequence database can aid in the identification of unknown proteins using tandem mass spectrometry.

Thus, there also exists a need in the art for polypeptides suitable for use in peptide fragmentation studies, preferably, polypeptides that have altered expression in irritable bowel diseases, for use in molecular weight measurements, and for use in protein sequencing using tandem mass spectrometry.

SUMMARY OF THE INVENTION

Using the T84 model, it was determined that certain polypeptides have altered (up-regulated or down-regulated) expression patterns in response to INF-gamma. Such molecules may have a role in gut barrier function and IBD and may be useful as potential therapeutic agents in the treatment of IBD and other gut pathologies. The invention aids in fulfilling these needs in the art by providing isolated nucleic acids and polypeptides encoded by these nucleic acids that have altered expression characteristics in the T84 gut barrier model. Particular embodiments of the invention are directed to isolated nucleic acid molecules comprising the DNA sequences of SEQ ID NOs:1-23 and isolated nucleic acid molecules encoding the amino acid sequences of SEQ ID NOs:24-33, as well as nucleic acid molecules complementary to these sequences. Both single-stranded and double-stranded RNA and DNA nucleic acid molecules are encompassed by the invention, as well as nucleic acid molecules that hybridize to a denatured, double-stranded DNA comprising all or a portion of SEQ ID NOs:1-23 and/or the DNA that encodes the amino acid sequences of SEQ ID NOs:24-33. Also encompassed are isolated nucleic acid molecules that are derived by *in vitro* mutagenesis from nucleic acid molecules comprising sequences of SEQ ID

NOs:1-23, that are degenerate from nucleic acid molecules comprising sequences of SEQ ID NOs:1-23, and that are allelic variants of DNA of the invention. The invention also encompasses recombinant vectors that direct the expression of these nucleic acid molecules and host cells transformed or transfected with these vectors.

5 In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%, more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO:1-23 or a polynucleotide which is
10 hybridizable to SEQ ID NO:1-23;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 1-23 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO:1-23;

(c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO:
15 1-23 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO:1-23;

(d) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 1-23, having biological activity;

(e) a polynucleotide which is a variant of SEQ ID NO:1-23;

20 (f) a polynucleotide which is an allelic variant of SEQ ID NO:1-23;

(g) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 1-23;

(h) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not
25 hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%, more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected
30 from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO:1, 7, 8, 10, 14-17, 19, 21 or a polynucleotide which is hybridizable to SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21;

(b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21;

5 (c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21;

(d) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21, having biological activity;

(e) a polynucleotide which is a variant of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21;

10 (f) a polynucleotide which is an allelic variant of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21;

(g) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 1, 7, 8, 10, 14-17, 19, 21, 7, 8, 10, 14-17, 19, 21;

15 (h) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

In another embodiment, the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 80%, preferable 85%,
20 more preferably 90%, optimally 95%, identical to a sequence of a polynucleotide selected from the group consisting of:

(a) a polynucleotide fragment of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23 or a polynucleotide which is hybridizable to SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23;

25 (b) a polynucleotide encoding a polypeptide fragment of a translation of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23 or a polypeptide fragment encoded by the cDNA sequence which is hybridizable to SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23;

(c) a polynucleotide encoding a polypeptide epitope of a translation of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23 or a polypeptide epitope encoded by a cDNA sequence which is hybridizable to SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23;

30 (d) a polynucleotide encoding a polypeptide of a translation of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23, having biological activity;

(e) a polynucleotide which is a variant of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23;

(f) a polynucleotide which is an allelic variant of SEQ ID NO:2-6, 9, 11-13, 18, 20, 22, 23;

(g) a polynucleotide which encodes a species homologue of a translation of SEQ ID NO: 2-6, 9, 11-13, 18, 20, 22, 23;

5 (h) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(g), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

10 In one embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a secreted protein. In preferred embodiments, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a polypeptide chosen from the group consisting of:

(a) a polypeptide having the polypeptide sequence identified as a translation of SEQ ID NO: 1-23;

15 (b) a polypeptide having the polypeptide sequence of SEQ ID NO: 24-33; and

(c) a polypeptide encoded by the cDNA which is hybridizable to SEQ ID NO:1-23.

Typically, the isolated nucleic acid molecule comprises the entire nucleotide sequence of SEQ ID NO:1-23 or a cDNA sequence which is hybridizable to SEQ ID NO:1-23.

20 Typically the isolated nucleic acid molecule comprises sequential nucleotide deletions from portions of the nucleotide sequence encoding either the C-terminus or the N-terminus of the polypeptide.

In other aspects, the invention provides a recombinant vector comprising the isolated nucleic acid molecule and a method of making a recombinant host cell comprising the isolated nucleic acid molecule and the recombinant host cell produced by such a method.

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In other preferred embodiments, the invention provides an isolated polypeptide having an amino acid sequence at least 80%, preferably at least 85%, more preferably at least 90% identical to the sequence of a polypeptide selected from the group consisting of:

30 (a) a polypeptide fragment of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

(b) a polypeptide having the sequence of SEQ ID NO: 24-33;

(c) a polypeptide domain of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

(d) a polypeptide epitope of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

(e) a secreted form of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

5 (f) a full length protein of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

(g) a variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23;

(h) an allelic variant of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23; and

10 (i) a species homologue of a polypeptide encoded by a polynucleotide of SEQ ID NO: 1-23.

In some embodiments, the full length polypeptide comprises sequential amino acid deletions from the C-terminus. In other embodiments, the mature polypeptide comprises sequential amino acid deletions from the C-terminus. Alternatively, the full length polypeptide can comprise sequential amino acid deletions from the N-terminus or the mature polypeptide can comprise sequential amino acid deletions from the N-terminus.

In other aspects, the invention provides an isolated antibody that binds specifically to the isolated polypeptide, a recombinant host cell that expresses the isolated polypeptide, and a method of making an isolated polypeptide comprising culturing the recombinant host cell under conditions such that said polypeptide is expressed; and recovering said polypeptide.

The invention in another embodiment is a method for preventing, treating, or ameliorating irritable bowel disorders, comprising administering to a mammalian subject a therapeutically effective amount of the isolated polypeptide or the isolated nucleic acid molecule. In another embodiment, the invention provides a method of diagnosing an irritable bowel disease or a susceptibility to irritable bowel disease in a subject comprising: determining the presence or absence of a mutation in the isolated nucleic acid molecule, and diagnosing an irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or absence of said mutation. Alternatively, irritable bowel disease or a susceptibility to irritable bowel disease in a subject can be diagnosed by a method comprising: determining the presence or amount of expression of the polypeptide in a

biological sample; and diagnosing irritable bowel disease or a susceptibility to irritable bowel disease based on the presence or amount of expression of the polypeptide.

Typically, a binding partner of the polypeptide is identified by contacting the polypeptide with a binding partner; and determining whether the binding partner affects a physical property or an activity of the polypeptide. Typically activity in a biological assay of a secreted polypeptide is identified by expressing the polynucleotide of SEQ ID NO:1-23 in a cell; isolating the supernatant; detecting an activity in a biological assay; and identifying the polypeptide in the supernatant having the activity.

10 In addition, the invention encompasses methods of using the nucleic acids noted above to identify nucleic acids encoding proteins homologous to SEQ ID NOs:24-33; to identify human chromosomes that contain the nucleotide sequences of the invention; to map genes near the nucleotide sequences of the invention on human chromosomes; and to identify genes associated with certain diseases, syndromes, or other human conditions associated with human chromosomes containing sequences of the invention.

For example, four of the nucleotide sequences of the invention, IMX4 and IMX56, IMX21, and IMX44 are located on chromosomes 22, 22, 7, and 19, respectively (IMX4 and IMX56 are both located on chromosome 22). Thus, the above-named nucleotide sequences (IMX4, IMX56, IMX21, and IMX44) can be used to identify human chromosome numbers 22, 7, and 19; to map genes on human chromosome numbers 22, 7, and 19; and to identify genes associated with certain diseases, syndromes, or other human conditions associated with human chromosome numbers 22, 7, and 19.

The invention also encompasses the use of sense or antisense oligonucleotides from the nucleic acids of SEQ ID NOs:1-23 to inhibit the expression of the polynucleotides encoded by the nucleotide sequences of the invention.

The invention also encompasses isolated polypeptides and fragments thereof encoded by these nucleic acid molecules including soluble polypeptide portions of SEQ ID NOs:24-33. The invention further encompasses methods for the production of these polypeptides, including culturing a host cell under conditions promoting expression and recovering the polypeptide from the culture medium if it is secreted or from cultured cells if it is not secreted. Especially, the expression of these polypeptides in bacteria, yeast, plant, insect, and animal cells is encompassed by the invention.

In addition, the invention includes assays utilizing these polypeptides, to screen for potential inhibitors of activity associated with polypeptide counter-structure molecules, and methods of using these polypeptides as therapeutic agents for the treatment of diseases mediated by polypeptide counter-structure molecules. Further, methods of using these polypeptides in the design of inhibitors thereof are also an aspect of the invention.

The invention further includes a method for using these polypeptides as molecular weight markers that allow the estimation of the molecular weight of a protein or a fragmented protein, as well as a method for the visualization of the molecular weight markers of the invention thereof using electrophoresis. The invention further encompasses methods for using the polypeptides of the invention as markers for determining the isoelectric point of an unknown protein, as well as controls for establishing the extent of fragmentation of a protein.

Further encompassed by this invention are kits to aid in these determinations.

Further encompassed by this invention is the use of the IMX nucleic acid sequences, predicted amino acid sequences of the polypeptide or fragments thereof, or a combination of the predicted amino acid sequences of the polypeptides and fragments thereof for use in searching an electronic database to aid in the identification of sample nucleic acids and/or proteins.

The invention also encompasses IMX polypeptides and the use of these polypeptides as research reagents to further study gut epithelial barrier function and regulation and therapeutic reagents to treat inflammatory bowel disease and other gut pathologies.

Isolated polyclonal or monoclonal antibodies that bind to these polypeptides are also encompassed by the invention, in addition the use of these antibodies to aid in purifying IMX polypeptides.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 presents the nucleotide sequence of IMX 4 (SEQ ID NO: 1).

Figure 2 presents the nucleotide sequence of IMX 10 (SEQ ID NO: 14).

Figure 3 presents the nucleotide sequence of IMX 21 (SEQ ID NO: 15).

Figure 4 presents the nucleotide sequence of IMX 28 (SEQ ID NO: 16).

Figure 5 presents the nucleotide sequence of IMX 32 (SEQ ID NO: 17).

Figure 6 presents the nucleotide sequence of IMX 39 (SEQ ID NO: 19).

Figure 7 presents the nucleotide sequence of IMX 40 (SEQ ID NO: 7).

Figure 8 presents the nucleotide sequence of IMX 42 (SEQ ID NO: 8).
Figure 9 presents the nucleotide sequence of IMX 44 (SEQ ID NO: 21).
Figure 10 presents the nucleotide sequence of IMX 56 (SEQ ID NO: 10).
Figure 11 presents the amino acid sequence of IMX 4 (SEQ ID NO: 24).
5 Figure 12 presents the amino acid sequence of IMX 10 (SEQ ID NO: 25).
Figure 13 presents the amino acid sequence of IMX 21 (SEQ ID NO: 26).
Figure 14 presents the amino acid sequence of IMX 28 (SEQ ID NO: 27).
Figure 15 presents the amino acid sequence of IMX 32 (SEQ ID NO: 28).
Figure 16 presents the amino acid sequence of IMX 39 (SEQ ID NO: 29).
10 Figure 17 presents the amino acid sequence of IMX 40 (SEQ ID NO: 30).
Figure 18 presents the amino acid sequence of IMX 42 (SEQ ID NO: 31).
Figure 19 presents the amino acid sequence of IMX 44 (SEQ ID NO: 32).
Figure 20 presents the amino acid sequence of IMX 56 (SEQ ID NO: 33).
Figure 21 presents the nucleotide sequence of the 5' end of the clone (SEQ ID
15 NO:13) matched to part of the human ApoL gene (AF019225).

Figure 22 presents comparison of the 5' end of the clone suggests it represents an alternative splice product to reported ApoL, i.e., bases 1-168 match to 2 exons on PAC carrying the ApoL gene but are not included in the reported complete cDNA.

DETAILED DESCRIPTION OF THE INVENTIONDefinitions

5 The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still
10 be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily
15 containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

As used herein, a "polynucleotide" refers to a molecule having a nucleic acid
20 sequence contained in SEQ ID NO:1-23. For example, the polynucleotide can contain all or part of the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the
25 translated amino acid sequence generated from the polynucleotide as broadly defined.

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:1-23, or the complement thereof, or the cDNA. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50%
30 formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH₂PO₄; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

Of course, a polynucleotide which hybridizes only to polyA⁺ sequences (such as any 3' terminal polyA⁺ tract of a cDNA shown in the sequence listing), or to a complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of

modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formulation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS – STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990); Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that

of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.).

The translated amino acid sequence, beginning with the methionine, is identified although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by the translation of these alternative open reading frames are specifically contemplated by the present invention.

SEQ ID NO:1-23 and the translations of SEQ ID NO: 1-23 as well as SEQ ID NO:24-33 are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from the translations of SEQ ID NO:1-23 may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified.

Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

The present invention also relates to the genes corresponding to SEQ ID NO:1-23, and translations of SEQ ID NO:1-23. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

Also provided in the present invention are species homologues. Species homologues may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for the desired homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood
5 in the art.

The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below). It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple
10 histidine residues, or an additional sequence for stability during recombinant production.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the
15 invention also can be purified from natural or recombinant sources using antibodies of the invention raised against the secreted protein in methods which are well known in the art.

Signal Sequences

Methods for predicting whether a protein has a signal sequence, as well as the
20 cleavage point for that sequence, are available. For instance, the method of McGeoch, Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 indicates the amino
25 terminus of the secreted protein. Therefore, from a deduced amino acid sequence, a signal sequence and mature sequence can be identified.

Polynucleotide and Polypeptide Variants

"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide
30 or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

Further embodiments of the present invention include polynucleotides having at least 80% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to a sequence contained in SEQ ID NO:1-23. Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the polynucleotides having at least 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity will encode a polypeptide identical to an amino acid sequence contained in the translations of SEQ ID NO:1-23.

Similarly, by a polypeptide having an amino acid sequence having at least, for example, 95% "identity" to a reference polypeptide, is intended that the amino acid sequence of the polypeptide is identical to the reference polypeptide except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the total length of the reference polypeptide. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in the reference sequence may be deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

Further embodiments of the present invention include polypeptides having at least 80% identity, more preferably at least 85% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to an amino acid sequence contained in translations of SEQ ID NO: 1-23. Preferably, the above polypeptides should exhibit at least one biological activity of the protein.

In a preferred embodiment, polypeptides of the present invention include polypeptides having at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98%, or 99% similarity to an amino acid sequence contained in translations of SEQ ID NO:1-23 as well as the amino acid sequences of SEQ ID NO:24-33.

The variants may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the

encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 (1993) reported variant KGF proteins having heparin binding activity even after deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)

Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1 a. They used random mutagenesis to generate over 3,500 individual IL-1 a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." (See Gayle et al., (1993), Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion

variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by
5 routine methods described herein and otherwise known in the art.

Thus, the invention further includes polypeptide variants which show substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid
10 substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change. As the authors state, these two strategies, natural selection and genetic engineering, have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be
15 permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr;
20 replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

Besides conservative amino acid substitution, variants of the present invention
25 include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv)
30 fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity.

5 (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

Polynucleotide and Polypeptide Fragments

10 In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in that shown in SEQ ID NO:1-23. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or
15 more contiguous bases from the cDNA sequence contained in that shown in SEQ ID NO:1-23. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, and more nucleotides) are preferred.

Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-
20 230, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, and so forth, to the end of SEQ ID NO:1-23. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity.

In the present invention, a "polypeptide fragment" refers to a short amino acid
25 sequence contained in the translations of SEQ ID NO:1-23 as well as SEQ ID NO:24-33. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, and so forth, to the end of the
30 coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50 or 60, amino acids in length. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of the translations of SEQ ID NO:1-23 as well as SEQ ID NO:24-33 falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

Epitopes & Antibodies

In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998-4002 (1983).)

Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

5 In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

10 Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least
15 about 25 amino acids), without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant
20 to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')₂ fragments) which are capable of specifically binding to protein. Fab and F(ab')₂ fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred, as well as the products of a
25 FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

Fusion Proteins

Any polypeptide of the present invention can be used to generate fusion proteins.
30 For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals,

the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., *Nature* 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., *J. Biochem.* 270:3958-3964 (1995).)

Similarly, EP-A-0 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. (EP-A 0 232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify

antagonists of hIL-5. (See, D. Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. Chem. 270:9459-9471 (1995).)

Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a peptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QLAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).)

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

15 **Vectors, Host Cells, and Protein Production**

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques.

The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

Nucleic Acid Molecules and Polypeptides of the Present Invention

The nucleic acid molecules encompassed in the invention comprise nucleotide sequences SEQ ID NO: 1-23.

25 The amino acid sequences of the polypeptides encoded by the nucleotide sequences of the invention are given in SEQ ID NO:24-33.

The discovery of the nucleic acids of the invention enables the construction of expression vectors comprising nucleic acid sequences encoding polypeptides; host cells transfected or transformed with the expression vectors; isolated and purified biologically active polypeptides and fragments thereof; the use of the nucleic acids or oligonucleotides thereof as probes to identify nucleic acids encoding proteins having amino acid sequences homologous to SEQ ID NOs: 24-33; the use of the nucleic acids or oligonucleotides thereof to identify human chromosomes, for example, 22, 7, and 19; the use of the nucleic acids or

oligonucleotides thereof to map genes on human chromosomes. for example, 22, 7, and 19; the use of the nucleic acids or oligonucleotides thereof to identify genes associated with certain diseases, syndromes or other human conditions associated with human chromosomes such as 22, 7, and 19; the use of single-stranded sense or antisense oligonucleotides from the nucleic acids to inhibit expression of polynucleotides encoded by the IMX sequences; the use of such polypeptides and soluble fragments as molecular weight markers; the use of such polypeptides and fragmented peptides as controls for peptide fragmentation, and kits comprising these reagents; the use of such polypeptides and fragments thereof to generate antibodies, and the use of antibodies to purify IMX polypeptides.

NUCLEIC ACID MOLECULES

In a particular embodiment, the invention relates to certain isolated nucleotide sequences that are free from contaminating endogenous material. A "nucleotide sequence" refers to a polynucleotide molecule in the form of a separate fragment or as a component of a larger nucleic acid construct. The nucleic acid molecule has been derived from DNA or RNA isolated at least once in substantially pure form and in a quantity or concentration enabling identification, manipulation, and recovery of its component nucleotide sequences by standard biochemical methods (such as those outlined in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY (1989)). Such sequences are preferably provided and/or constructed in the form of an open reading frame uninterrupted by internal non-translated sequences, or introns, that are typically present in eukaryotic genes. Sequences of non-translated DNA can be present 5' or 3' from an open reading frame, where the same do not interfere with manipulation or expression of the coding region.

Nucleic acid molecules of the invention include DNA in both single-stranded and double-stranded form, as well as the RNA complement thereof. DNA includes, for example, cDNA, genomic DNA, chemically synthesized DNA, DNA amplified by PCR, and combinations thereof. Genomic DNA may be isolated by conventional techniques, e.g., using the cDNA of SEQ ID NOs:1-23, or a suitable fragment thereof, as a probe.

The DNA molecules of the invention include full length genes as well as polynucleotides and fragments thereof. The full length gene may include the N-terminal signal peptide. Other embodiments include DNA encoding a soluble form, e.g., encoding the extracellular domain of the protein, either with or without the signal peptide.

The nucleic acids of the invention are preferentially derived from human sources, but the invention includes those derived from non-human species, as well.

Preferred Sequences

The particularly preferred nucleotide sequences of the invention are SEQ ID NOs: 1-23, as set forth above. cDNA clones having the nucleotide sequence of SEQ ID NOs:1-23 were isolated as described in Example 1. The sequences of amino acids encoded by the DNA of SEQ ID NOs:1-23 are shown in SEQ ID NOs:24-33.

Additional Sequences

Due to the known degeneracy of the genetic code, wherein more than one codon can encode the same amino acid, a DNA sequence can vary from that shown in SEQ ID NOs:1-23, and still encode a polypeptide having the amino acid sequence of SEQ ID NOs:24-33. Such variant DNA sequences can result from silent mutations (e.g., occurring during PCR amplification), or can be the product of deliberate mutagenesis of a native sequence.

The invention thus provides additional isolated DNA sequences encoding polypeptides of the invention, selected from: (a) DNA comprising the nucleotide sequences of SEQ ID NOs: 1-23; (b) DNA encoding the polypeptides of SEQ ID NOs:24-33; (c) DNA capable of hybridization to a DNA of (a) or (b) under conditions of moderate stringency and which encodes polypeptides of the invention; (d) DNA capable of hybridization to a DNA of (a) or (b) under conditions of high stringency and which encodes polypeptides of the invention, and (e) DNA which is degenerate, as a result of the genetic code, to a DNA defined in (a), (b), (c), or (d) and which encode polypeptides of the invention. Of course, polypeptides encoded by such DNA sequences are encompassed by the invention.

As used herein, conditions of moderate stringency can be readily determined by those having ordinary skill in the art based on, for example, the length of the DNA. The basic conditions are set forth by Sambrook et al. *Molecular Cloning: A Laboratory Manual*, 2 ed. Vol. 1, pp. 1.101-234, Cold Spring Harbor Laboratory Press, (1989), and include use of a prewashing solution for the nitrocellulose filters 5X SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0), hybridization conditions of about 50% formamide, 6X SSC at about 42°C (or other similar hybridization solution, such as Stark's solution, in about 50% formamide at about 42°C), and washing conditions of about 60°C, 0.5X SSC, 0.1% SDS. Conditions of high stringency can also be readily determined by the skilled artisan based on,

for example, the length of the DNA. Generally, such conditions are defined as hybridization conditions as above, and with washing at approximately 68°C, 0.2X SSC, 0.1% SDS. The skilled artisan will recognize that the temperature and wash solution salt concentration can be adjusted as necessary according to factors such as the length of the probe.

5 Also included as an embodiment of the invention is DNA encoding polypeptide fragments and polypeptides comprising inactivated N-glycosylation site(s), inactivated protease processing site(s), or conservative amino acid substitution(s), as described below.

In another embodiment, the nucleic acid molecules of the invention also comprise nucleotide sequences that are at least 80% identical to a native sequence. Also
10 contemplated are embodiments in which a nucleic acid molecule comprises a sequence that is at least 90% identical, at least 95% identical, at least 98% identical, at least 99% identical, or at least 99.9% identical to a native sequence.

The percent identity may be determined by visual inspection and mathematical calculation. Alternatively, the percent identity of two nucleic acid sequences can be
15 determined by comparing sequence information using the GAP computer program, version 6.0 described by Devereux et al. (*Nucl. Acids Res.* 12:387, 1984) and available from the University of Wisconsin Genetics Computer Group (UWGCG). The preferred default parameters for the GAP program include: (1) a unary comparison matrix (containing a value of 1 for identities and 0 for non-identities) for nucleotides, and the weighted
20 comparison matrix of Gribskov and Burgess, *Nucl. Acids Res.* 14:6745, 1986, as described by Schwartz and Dayhoff, eds., *Atlas of Protein Sequence and Structure*, National Biomedical Research Foundation, pp. 353-358, 1979; (2) a penalty of 3.0 for each gap and an additional 0.10 penalty for each symbol in each gap; and (3) no penalty for end gaps. Other programs used by one skilled in the art of sequence comparison may also be used.

25 The invention provides isolated nucleic acids useful in the production of polypeptides. Such polypeptides may be prepared by any of a number of conventional techniques. A DNA sequence encoding an IMX polypeptide or a desired fragment thereof may be subcloned into an expression vector for production of the polypeptide or fragment. The DNA sequence advantageously is fused to a sequence encoding a suitable leader or
30 signal peptide. Alternatively, the desired fragment may be chemically synthesized using known techniques. DNA fragments also may be produced by restriction endonuclease digestion of a full length cloned DNA sequence, and isolated by electrophoresis on agarose gels. If necessary, oligonucleotides that reconstruct the 5' or 3' terminus to a desired point

may be ligated to a DNA fragment generated by restriction enzyme digestion. Such oligonucleotides may additionally contain a restriction endonuclease cleavage site upstream of the desired coding sequence, and position an initiation codon (ATG) at the N-terminus of the coding sequence.

5 The well-known polymerase chain reaction (PCR) procedure also may be employed to isolate and amplify a DNA sequence encoding a desired protein fragment.

Oligonucleotides that define the desired termini of the DNA fragment are employed as 5' and 3' primers. The oligonucleotides may additionally contain recognition sites for

restriction endonucleases, to facilitate insertion of the amplified DNA fragment into an

10 expression vector. PCR techniques are described in Saiki et al., *Science* 239:487 (1988);

Recombinant DNA Methodology; Wu et al., eds., Academic Press, Inc., San Diego (1989),

pp. 189-196; and *PCR Protocols: A Guide to Methods and Applications*, Innis et al., eds.,

Academic Press, Inc. (1990).

15 POLYPEPTIDES AND FRAGMENTS THEREOF

 The invention encompasses polypeptides and fragments thereof in various forms, including those that are naturally occurring or produced through various techniques such as procedures involving recombinant DNA technology. Such forms include, but are not limited to, derivatives, variants, and oligomers, as well as fusion proteins or fragments

20 thereof.

Polypeptides and Fragments Thereof

 The polypeptides of the invention include full length proteins encoded by the nucleic acid sequences set forth above. Particularly preferred polypeptides comprise the amino acid

25 sequences of SEQ ID NOs:24-33.

 The polypeptides of the invention may include an N-terminal hydrophobic region that functions as a signal peptide, and may contain an extracellular domain, and may also contain a transmembrane region and a C-terminal cytoplasmic domain as well as a spacer region. Computer analysis may be used to predict the location of the signal peptide.

30 For example, the isolated polypeptides of SEQ ID NO:16 (IMX 28) and SEQ ID NO:21 (IMX 44) include an N-terminal hydrophobic region that functions as a signal peptide. Computer analysis predicts that the signal peptide corresponds to residues 1 to 17 of SEQ ID NO:16, and cleavage of the signal peptide of SEQ ID NO:16 results in a mature

protein comprising amino acids 18 to 372. The signal peptide of EMX 44 corresponds to residues 1 to 37 of SEQ ID NO:21. The next most likely computer-predicted signal peptide cleavage sites (in descending order) would occur after amino acids 36, 26 and 27 of SEQ ID NO:21. Cleavage of the signal peptide at position 37 thus would yield a mature protein
5 comprising amino acids 38 through 261 of SEQ ID NO:21.

The skilled artisan will recognize that the above-described boundaries of such regions of the polypeptide are approximate. To illustrate, the boundaries of the mature protein (which may be predicted by using computer programs available for that purpose) may differ from those described above.

10 The polypeptides of the invention may be membrane bound or they may be secreted and, thus, soluble. Soluble polypeptides are capable of being secreted from the cells in which they are expressed. In general, soluble polypeptides may be identified (and distinguished from non-soluble membrane-bound counterparts) by separating intact cells which express the desired polypeptide from the culture medium, *e.g.*, by centrifugation, and
15 assaying the medium (supernatant) for the presence of the desired polypeptide. The presence of polypeptide in the medium indicates that the polypeptide was secreted from the cells and thus is a soluble form of the protein.

In one embodiment, the soluble polypeptides and fragments thereof comprise all or part of the extracellular domain, but lack the transmembrane region that would cause
20 retention of the polypeptide on a cell membrane. A soluble polypeptide may include the cytoplasmic domain, or a portion thereof, as long as the polypeptide is secreted from the cell in which it is produced.

In general, the use of soluble forms is advantageous for certain applications. Purification of the polypeptides from recombinant host cells is facilitated, since the soluble
25 polypeptides are secreted from the cells. Further, soluble polypeptides are generally more suitable for intravenous administration.

The invention also provides polypeptides and fragments of the extracellular domain that retain a desired biological activity. Particular embodiments are directed to polypeptide fragments that retain the ability to bind the native cognate, substrate, or counter-structure
30 ("binding partner"). Such a fragment may be a soluble polypeptide, as described above. In another embodiment, the polypeptides and fragments advantageously include regions that are conserved in the family as described above.

Also provided herein are polypeptide fragments comprising at least 20, or at least 30, contiguous amino acids of the sequence of SEQ ID NOs:24-33. Fragments derived from the cytoplasmic domain find use in studies of signal transduction, and in regulating cellular processes associated with transduction of biological signals. Polypeptide fragments also
5 may be employed as immunogens, in generating antibodies.

Variants

Naturally occurring variants as well as derived variants of the polypeptides and fragments are provided herein.

10 Variants may exhibit amino acid sequences that are at least 80% identical. Also contemplated are embodiments in which a polypeptide or fragment comprises an amino acid sequence that is at least 90% identical, at least 95% identical, at least 98% identical, at least 99% identical, or at least 99.9% identical to the preferred polypeptide or fragment thereof. Percent identity may be determined by visual inspection and mathematical calculation.

15 Alternatively, the percent identity of two protein sequences can be determined by comparing sequence information using the GAP computer program, based on the algorithm of Needleman and Wunsch (J. Mol. Bio. 48:443, 1970) and available from the University of Wisconsin Genetics Computer Group (UWGCG). The preferred default parameters for the GAP program include: (1) a scoring matrix, blosum62, as described by Henikoff and
20 Henikoff (Proc. Natl. Acad. Sci. USA 89:10915, 1992); (2) a gap weight of 12; (3) a gap length weight of 4; and (4) no penalty for end gaps. Other programs used by one skilled in the art of sequence comparison may also be used.

The variants of the invention include, for example, those that result from alternate mRNA splicing events or from proteolytic cleavage. Alternate splicing of mRNA may, for
25 example, yield a truncated but biologically active protein, such as a naturally occurring soluble form of the protein. Variations attributable to proteolysis include, for example, differences in the N- or C-termini upon expression in different types of host cells, due to proteolytic removal of one or more terminal amino acids from the protein (generally from 1-5 terminal amino acids). Proteins in which differences in amino acid sequence are
30 attributable to genetic polymorphism (allelic variation among individuals producing the protein) are also contemplated herein.

Additional variants within the scope of the invention include polypeptides that may be modified to create derivatives thereof by forming covalent or aggregative conjugates with

other chemical moieties, such as glycosyl groups, lipids, phosphate, acetyl groups and the like. Covalent derivatives may be prepared by linking the chemical moieties to functional groups on amino acid side chains or at the N-terminus or C-terminus of a polypeptide. Conjugates comprising diagnostic (detectable) or therapeutic agents attached thereto are contemplated herein, as discussed in more detail below.

Other derivatives include covalent or aggregative conjugates of the polypeptides with other proteins or polypeptides, such as by synthesis in recombinant culture as N-terminal or C-terminal fusions. Examples of fusion proteins are discussed below in connection with oligomers. Further, fusion proteins can comprise peptides added to facilitate purification and identification. Such peptides include, for example, poly-His or the antigenic identification peptides described in U.S. Patent No. 5,011,912 and in Hopp et al., *Bio/Technology* 6:1204, 1988. One such peptide is the FLAG[®] peptide, Asp-Tyr-Lys-Asp-Asp-Asp-Lys, which is highly antigenic and provides an epitope reversibly bound by a specific monoclonal antibody, enabling rapid assay and facile purification of expressed recombinant protein. A murine hybridoma designated 4E11 produces a monoclonal antibody that binds the FLAG[®] peptide in the presence of certain divalent metal cations, as described in U.S. Patent 5,011,912, hereby incorporated by reference. The 4E11 hybridoma cell line has been deposited with the American Type Culture Collection under accession no. HB 9259. Monoclonal antibodies that bind the FLAG[®] peptide are available from Eastman Kodak Co., Scientific Imaging Systems Division, New Haven, Connecticut.

Among the polypeptides provided herein are variants of native polypeptides that retain the native biological activity or the substantial equivalent thereof. One example is a variant that binds with essentially the same binding affinity as does the native form. Binding affinity can be measured by conventional procedures, *e.g.*, as described in U.S. Patent No. 5,512,457 and as set forth below.

Variants include polypeptides that are substantially homologous to the native form, but which have an amino acid sequence different from that of the native form because of one or more deletions, insertions or substitutions. Particular embodiments include, but are not limited to, polypeptides that comprise from one to ten deletions, insertions or substitutions of amino acid residues, when compared to a native sequence.

A given amino acid may be replaced, for example, by a residue having similar physiochemical characteristics. Examples of such conservative substitutions include substitution of one aliphatic residue for another, such as Ile, Val, Leu, or Ala for one

another: substitutions of one polar residue for another, such as between Lys and Arg, Glu and Asp, or Gln and Asn; or substitutions of one aromatic residue for another, such as Phe, Trp, or Tyr for one another. Other conservative substitutions, *e.g.*, involving substitutions of entire regions having similar hydrophobicity characteristics, are well known.

5 Similarly, the DNAs of the invention include variants that differ from a native DNA sequence because of one or more deletions, insertions or substitutions, but that encode a biologically active polypeptide.

The invention further includes polypeptides of the invention with or without associated native-pattern glycosylation. Polypeptides expressed in yeast or mammalian
10 expression systems (*e.g.*, COS-1 or COS-7 cells) can be similar to or significantly different from a native polypeptide in molecular weight and glycosylation pattern, depending upon the choice of expression system. Expression of polypeptides of the invention in bacterial expression systems, such as *E. coli*, provides non-glycosylated molecules. Further, a given preparation may include multiple differentially glycosylated species of the protein. Glycosyl
15 groups can be removed through conventional methods, in particular those utilizing glycopeptidase. In general, glycosylated polypeptides of the invention can be incubated with a molar excess of glycopeptidase (Boehringer Mannheim).

Correspondingly, similar DNA constructs that encode various additions or substitutions of amino acid residues or sequences, or deletions of terminal or internal
20 residues or sequences are encompassed by the invention. For example, N-glycosylation sites in the polypeptide extracellular domain can be modified to preclude glycosylation, allowing expression of a reduced carbohydrate analog in mammalian and yeast expression systems. N-glycosylation sites in eukaryotic polypeptides are characterized by an amino acid triplet Asn-X-Y, wherein X is any amino acid except Pro and Y is Ser or Thr.

25 Appropriate substitutions, additions, or deletions to the nucleotide sequence encoding these triplets will result in prevention of attachment of carbohydrate residues at the Asn side chain. Alteration of a single nucleotide, chosen so that Asn is replaced by a different amino acid, for example, is sufficient to inactivate an N-glycosylation site. Alternatively, the Ser or Thr can be replaced with another amino acid, such as Ala. Known procedures for
30 inactivating N-glycosylation sites in proteins include those described in U.S. Patent 5,071,972 and EP 276,846, hereby incorporated by reference.

In another example of variants, sequences encoding Cys residues that are not essential for biological activity can be altered to cause the Cys residues to be deleted or

replaced with other amino acids, preventing formation of incorrect intramolecular disulfide bridges upon folding or renaturation.

Other variants are prepared by modification of adjacent dibasic amino acid residues, to enhance expression in yeast systems in which KEX2 protease activity is present. EP 212,914 discloses the use of site-specific mutagenesis to inactivate KEX2 protease processing sites in a protein. KEX2 protease processing sites are inactivated by deleting, adding or substituting residues to alter Arg-Arg, Arg-Lys, and Lys-Arg pairs to eliminate the occurrence of these adjacent basic residues. Lys-Lys pairings are considerably less susceptible to KEX2 cleavage, and conversion of Arg-Lys or Lys-Arg to Lys-Lys represents a conservative and preferred approach to inactivating KEX2 sites.

Oligomers

Encompassed by the invention are oligomers or fusion proteins that contain IMX polypeptides. Such oligomers may be in the form of covalently-linked or non-covalently-linked multimers, including dimers, trimers, or higher oligomers. As noted above, preferred polypeptides are soluble and thus these oligomers may comprise soluble polypeptides. In one aspect of the invention, the oligomers maintain the binding ability of the polypeptide components and provide therefor, bivalent, trivalent, etc., binding sites.

One embodiment of the invention is directed to oligomers comprising multiple polypeptides joined *via* covalent or non-covalent interactions between peptide moieties fused to the polypeptides. Such peptides may be peptide linkers (spacers), or peptides that have the property of promoting oligomerization. Leucine zippers and certain polypeptides derived from antibodies are among the peptides that can promote oligomerization of the polypeptides attached thereto, as described in more detail below.

Immunoglobulin-based Oligomers

As one alternative, an oligomer is prepared using polypeptides derived from immunoglobulins. Preparation of fusion proteins comprising certain heterologous polypeptides fused to various portions of antibody-derived polypeptides (including the Fc domain) has been described, *e.g.*, by Ashkenazi et al. (*PNAS USA* 88:10535, 1991); Byrn et al. (*Nature* 344:677, 1990); and Hollenbaugh and Aruffo ("Construction of Immunoglobulin Fusion Proteins", in *Current Protocols in Immunology*, Suppl. 4, pages 10.19.1 - 10.19.11, 1992).

One embodiment of the present invention is directed to a dimer comprising two fusion proteins created by fusing a polypeptide of the invention to an Fc polypeptide derived from an antibody. A gene fusion encoding the polypeptide/Fc fusion protein is inserted into an appropriate expression vector. Polypeptide/Fc fusion proteins are expressed in host cells transformed with the recombinant expression vector, and allowed to assemble much like antibody molecules, whereupon interchain disulfide bonds form between the Fc moieties to yield divalent molecules.

The term "Fc polypeptide" as used herein includes native and mutein forms of polypeptides comprising from the Fc region of an antibody. Truncated forms of such polypeptides containing the hinge region that promotes dimerization are also included. Preferred polypeptides comprise an Fc polypeptide derived from a human IgG1 antibody comprising any or all of the CH domains of the Fc region.

One suitable Fc polypeptide, described in PCT application WO 93/10151, hereby incorporated by reference, is a single chain polypeptide extending from the N-terminal hinge region to the native C-terminus of the Fc region of a human IgG1 antibody. Another useful Fc polypeptide is the Fc mutein described in U.S. Patent 5,457,035 and in Baum et al., (*EMBO J.* 13:3992-4001, 1994) incorporated herein by reference. The amino acid sequence of this mutein is identical to that of the native Fc sequence presented in WO 93/10151, except that amino acid 19 has been changed from Leu to Ala, amino acid 20 has been changed from Leu to Glu, and amino acid 22 has been changed from Gly to Ala. The mutein exhibits reduced affinity for Fc receptors.

The above-described fusion proteins comprising Fc moieties (and oligomers formed therefrom) offer the advantage of facile purification by affinity chromatography over Protein A or Protein G columns.

In other embodiments, the polypeptides of the invention may be substituted for the variable portion of an antibody heavy or light chain. If fusion proteins are made with both heavy and light chains of an antibody, it is possible to form an oligomer with as many as four soluble regions of the proteins of the invention.

Peptide-linker Based Oligomers

Alternatively, the oligomer is a fusion protein comprising multiple polypeptides, with or without peptide linkers (spacer peptides). Among the suitable peptide linkers are those described in U.S. Patents 4,751,180 and 4,935,233, which are hereby incorporated by reference. A DNA sequence encoding a desired peptide linker may be inserted between, and in the same reading frame as, the DNA sequences of the invention, using any suitable conventional technique. For example, a chemically synthesized oligonucleotide encoding the linker may be ligated between the sequences. In particular embodiments, a fusion protein comprises from two to four soluble IMX polypeptides, separated by peptide linkers.

Leucine-Zippers

Another method for preparing the oligomers of the invention involves use of a leucine zipper. Leucine zipper domains are peptides that promote oligomerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding proteins (Landschulz et al., *Science* 240:1759, 1988), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize.

Examples of leucine zipper domains suitable for producing soluble oligomeric proteins are described in PCT application WO 94/10308, and the leucine zipper derived from lung surfactant protein D (SPD) described in Hoppe et al. (*FEBS Letters* 344:191, 1994), hereby incorporated by reference. The use of a modified leucine zipper that allows for stable trimerization of a heterologous protein fused thereto is described in Fanslow et al. (*Semin. Immunol.* 6:267-278, 1994). Recombinant fusion proteins comprising a soluble polypeptide fused to a leucine zipper peptide are expressed in suitable host cells, and the soluble oligomer that forms is recovered from the culture supernatant.

In particular embodiments, leucine residues in a leucine zipper moiety are replaced by isoleucine residues. Such peptides comprising isoleucine may be referred to as "isoleucine zippers" but are encompassed by the term "leucine zippers" as employed herein.

PRODUCTION OF POLYPEPTIDES AND FRAGMENTS THEREOF

Expression, isolation and purification of the polypeptides and fragments of the invention may be accomplished by any suitable technique, including but not limited to the following:

Expression Systems

The present invention also provides recombinant cloning and expression vectors containing DNA, as well as host cell containing the recombinant vectors. Expression
5 vectors comprising DNA may be used to prepare the polypeptides or fragments of the invention encoded by the DNA. A method for producing polypeptides comprises culturing host cells transformed with a recombinant expression vector encoding the polypeptide, under conditions that promote expression of the polypeptide, then recovering the expressed polypeptides from the culture. The skilled artisan will recognize that the procedure for
10 purifying the expressed polypeptides will vary according to such factors as the type of host cells employed, and whether the polypeptide is membrane-bound or a soluble form that is secreted from the host cell.

Any suitable expression system may be employed. The vectors include a DNA encoding a polypeptide or fragment of the invention, operably linked to suitable
15 transcriptional or translational regulatory nucleotide sequences, such as those derived from a mammalian, microbial, viral, or insect gene. Examples of regulatory sequences include transcriptional promoters, operators, or enhancers, an mRNA ribosomal binding site, and appropriate sequences which control transcription and translation initiation and termination. Nucleotide sequences are operably linked when the regulatory sequence functionally relates
20 to the DNA sequence. Thus, a promoter nucleotide sequence is operably linked to a DNA sequence if the promoter nucleotide sequence controls the transcription of the DNA sequence. An origin of replication that confers the ability to replicate in the desired host cells, and a selection gene by which transformants are identified, are generally incorporated into the expression vector.

25 In addition, a sequence encoding an appropriate signal peptide (native or heterologous) can be incorporated into expression vectors. A DNA sequence for a signal peptide (secretory leader) may be fused in frame to the nucleic acid sequence of the invention so that the DNA is initially transcribed, and the mRNA translated, into a fusion protein comprising the signal peptide. A signal peptide that is functional in the intended
30 host cells promotes extracellular secretion of the polypeptide. The signal peptide is cleaved from the polypeptide upon secretion of polypeptide from the cell.

The skilled artisan will also recognize that the position(s) at which the signal peptide is cleaved may differ from that predicted by computer program, and may vary according to

such factors as the type of host cells employed in expressing a recombinant polypeptide. A protein preparation may include a mixture of protein molecules having different N-terminal amino acids, resulting from cleavage of the signal peptide at more than one site.

Suitable host cells for expression of polypeptides include prokaryotes, yeast or
5 higher eukaryotic cells. Mammalian or insect cells are generally preferred for use as host cells. Appropriate cloning and expression vectors for use with bacterial, fungal, yeast, and mammalian cellular hosts are described, for example, in Pouwels et al. *Cloning Vectors: A Laboratory Manual*, Elsevier, New York, (1985). Cell-free translation systems could also
10 be employed to produce polypeptides using RNAs derived from DNA constructs disclosed herein.

Prokaryotic Systems

Prokaryotes include gram-negative or gram-positive organisms. Suitable prokaryotic host cells for transformation include, for example, *E. coli*, *Bacillus subtilis*, *Salmonella*
15 *typhimurium*, and various other species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*. In a prokaryotic host cell, such as *E. coli*, a polypeptide may include an N-terminal methionine residue to facilitate expression of the recombinant polypeptide in the prokaryotic host cell. The N-terminal Met may be cleaved from the expressed recombinant polypeptide.

20 Expression vectors for use in prokaryotic host cells generally comprise one or more phenotypic selectable marker genes. A phenotypic selectable marker gene is, for example, a gene encoding a protein that confers antibiotic resistance or that supplies an autotrophic requirement. Examples of useful expression vectors for prokaryotic host cells include those derived from commercially available plasmids such as the cloning vector pBR322 (ATCC
25 37017). pBR322 contains genes for ampicillin and tetracycline resistance and thus provides simple means for identifying transformed cells. An appropriate promoter and a DNA sequence are inserted into the pBR322 vector. Other commercially available vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and pGEM1 (Promega Biotec, Madison, WI, USA).

30 Promoter sequences commonly used for recombinant prokaryotic host cell expression vectors include β -lactamase (penicillinase), lactose promoter system (Chang et al., *Nature* 275:615, 1978; and Goeddel et al., *Nature* 281:544, 1979), tryptophan (trp) promoter system (Goeddel et al., *Nucl. Acids Res.* 8:4057, 1980; and EP-A-36776) and tac

promoter (Maniatis. *Molecular Cloning: A Laboratory Manual*. Cold Spring Harbor Laboratory. p. 412, 1982). A particularly useful prokaryotic host cell expression system employs a phage 8P_L promoter and a cI857ts thermolabile repressor sequence. Plasmid vectors available from the American Type Culture Collection which incorporate derivatives
5 of the 8P_L promoter include plasmid pHUB2 (resident in *E. coli* strain JMB9, ATCC 37092) and pPLc28 (resident in *E. coli* RR1, ATCC 53082).

Yeast Systems

Alternatively, the polypeptides may be expressed in yeast host cells, preferably from
10 the *Saccharomyces* genus (e.g., *S. cerevisiae*). Other genera of yeast, such as *Pichia* or *Kluyveromyces*, may also be employed. Yeast vectors will often contain an origin of replication sequence from a 2 μ yeast plasmid, an autonomously replicating sequence (ARS), a promoter region, sequences for polyadenylation, sequences for transcription termination, and a selectable marker gene. Suitable promoter sequences for yeast vectors include, among
15 others, promoters for metallothionein, 3-phosphoglycerate kinase (Hitzeman et al., *J. Biol. Chem.* 255:2073, 1980) or other glycolytic enzymes (Hess et al., *J. Adv. Enzyme Reg.* 7:149, 1968; and Holland et al., *Biochem.* 17:4900, 1978), such as enolase, glyceraldehyde-3-phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phosphofructokinase, glucose-6-phosphate isomerase, 3-phosphoglycerate mutase, pyruvate kinase,
20 triosephosphate isomerase, phospho-glucose isomerase, and glucokinase. Other suitable vectors and promoters for use in yeast expression are further described in Hitzeman, EPA-73,657. Another alternative is the glucose-repressible ADH2 promoter described by Russell et al. (*J. Biol. Chem.* 258:2674, 1982) and Beier et al. (*Nature* 300:724, 1982). Shuttle vectors replicable in both yeast and *E. coli* may be constructed by inserting DNA sequences
25 from pBR322 for selection and replication in *E. coli* (Amp^r gene and origin of replication) into the above-described yeast vectors.

The yeast ∇ -factor leader sequence may be employed to direct secretion of the polypeptide. The ∇ -factor leader sequence is often inserted between the promoter sequence and the structural gene sequence. See, e.g., Kurjan et al., *Cell* 30:933, 1982 and Bitter et al.,
30 *Proc. Natl. Acad. Sci. USA* 81:5330, 1984. Other leader sequences suitable for facilitating secretion of recombinant polypeptides from yeast hosts are known to those of skill in the art. A leader sequence may be modified near its 3' end to contain one or more restriction sites. This will facilitate fusion of the leader sequence to the structural gene.

Yeast transformation protocols are known to those of skill in the art. One such protocol is described by Hinnen et al., *Proc. Natl. Acad. Sci. U.S.A* 75:1929, 1978. The Hinnen et al. protocol selects for Trp⁻ transformants in a selective medium, wherein the selective medium consists of 0.67% yeast nitrogen base, 0.5% casamino acids, 2% glucose, 10 mg/ml adenine and 20 mg/ml uracil.

Yeast host cells transformed by vectors containing an ADH2 promoter sequence may be grown for inducing expression in a "rich" medium. An example of a rich medium is one consisting of 1% yeast extract, 2% peptone, and 1% glucose supplemented with 80 mg/ml adenine and 80 mg/ml uracil. Derepression of the ADH2 promoter occurs when glucose is exhausted from the medium.

Mammalian or Insect Systems

Mammalian or insect host cell culture systems also may be employed to express recombinant polypeptides. Baculovirus systems for production of heterologous proteins in insect cells are reviewed by Luckow and Summers, *Bio/Technology* 6:47 (1988).

Established cell lines of mammalian origin also may be employed. Examples of suitable mammalian host cell lines include the COS-7 line of monkey kidney cells (ATCC CRL 1651) (Gluzman et al., *Cell* 23:175, 1981), L cells, C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells, HeLa cells, and BHK (ATCC CRL 10) cell lines, and the CV1/EBNA cell line derived from the African green monkey kidney cell line CV1 (ATCC CCL 70) as described by McMahan et al. (*EMBO J.* 10: 2821, 1991).

Established methods for introducing DNA into mammalian cells have been described (Kaufman, R.J., *Large Scale Mammalian Cell Culture*, 1990, pp. 15-69). Additional protocols using commercially available reagents, such as Lipofectamine lipid reagent (Gibco/BRL) or Lipofectamine-Plus lipid reagent, can be used to transfect cells (Felgner et al., *Proc. Natl. Acad. Sci. USA* 84:7413-7417, 1987). In addition, electroporation can be used to transfect mammalian cells using conventional procedures, such as those in Sambrook et al. (*Molecular Cloning: A Laboratory Manual*, 2 ed. Vol. 1-3, Cold Spring Harbor Laboratory Press, 1989). Selection of stable transformants can be performed using methods known in the art, such as, for example, resistance to cytotoxic drugs. Kaufman et al., *Meth. in Enzymology* 185:487-511, 1990, describes several selection schemes, such as dihydrofolate reductase (DHFR) resistance. A suitable host strain for DHFR selection can be CHO strain DX-B11, which is deficient in DHFR (Urlaub and

Chasin, *Proc. Natl. Acad. Sci. USA* 77:4216-4220, 1980). A plasmid expressing the DHFR cDNA can be introduced into strain DX-B11, and only cells that contain the plasmid can grow in the appropriate selective media. Other examples of selectable markers that can be incorporated into an expression vector include cDNAs conferring resistance to antibiotics, such as G418 and hygromycin B. Cells harboring the vector can be selected on the basis of resistance to these compounds.

Transcriptional and translational control sequences for mammalian host cell expression vectors can be excised from viral genomes. Commonly used promoter sequences and enhancer sequences are derived from polyoma virus, adenovirus 2, simian virus 40 (SV40), and human cytomegalovirus. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early and late promoter, enhancer, splice, and polyadenylation sites can be used to provide other genetic elements for expression of a structural gene sequence in a mammalian host cell. Viral early and late promoters are particularly useful because both are easily obtained from a viral genome as a fragment, which can also contain a viral origin of replication (Fiers et al., *Nature* 273:113, 1978; Kaufman, *Meth. in Enzymology*, 1990). Smaller or larger SV40 fragments can also be used, provided the approximately 250 bp sequence extending from the *Hind* III site toward the *Bgl* I site located in the SV40 viral origin of replication site is included.

Additional control sequences shown to improve expression of heterologous genes from mammalian expression vectors include such elements as the expression augmenting sequence element (EASE) derived from CHO cells (Morris et al., *Animal Cell Technology*, 1997, pp. 529-534 and PCT Application WO 97/25420) and the tripartite leader (TPL) and VA gene RNAs from Adenovirus 2 (Gingras et al., *J. Biol. Chem.* 257:13475-13491, 1982). The internal ribosome entry site (IRES) sequences of viral origin allows dicistronic mRNAs to be translated efficiently (Oh and Sarnow, *Current Opinion in Genetics and Development* 3:295-300, 1993; Ramesh et al., *Nucleic Acids Research* 24:2697-2700, 1996).

Expression of a heterologous cDNA as part of a dicistronic mRNA followed by the gene for a selectable marker (e.g. DHFR) has been shown to improve transfectability of the host and expression of the heterologous cDNA (Kaufman, *Meth. in Enzymology*, 1990). Exemplary expression vectors that employ dicistronic mRNAs are pTR-DC/GFP described by Mosser et al., *Biotechniques* 22:150-161, 1997, and p2A5I described by Morris et al., *Animal Cell Technology*, 1997, pp. 529-534.

A useful high expression vector, pCAVNOT, has been described by Mosley et al., *Cell* 59:335-348, 1989. Other expression vectors for use in mammalian host cells can be constructed as disclosed by Okayama and Berg (*Mol. Cell. Biol.* 3:280, 1983). A useful system for stable high level expression of mammalian cDNAs in C127 murine mammary epithelial cells can be constructed substantially as described by Cosman et al. (*Mol. Immunol.* 23:935, 1986). A useful high expression vector, PMLSV N1/N4, described by Cosman et al., *Nature* 312:768, 1984, has been deposited as ATCC 39890. Additional useful mammalian expression vectors are described in EP-A-0367566, and in WO 91 18982, incorporated by reference herein. In yet another alternative, the vectors can be derived from retroviruses.

Another useful expression vector, pFLAG[®], can be used. FLAG[®] technology is centered on the fusion of a low molecular weight (1kD), hydrophilic, FLAG[®] marker peptide to the N-terminus of a recombinant protein expressed by pFLAG[®] expression vectors.

Regarding signal peptides that may be employed, the native signal peptide may be replaced by a heterologous signal peptide or leader sequence, if desired. The choice of signal peptide or leader may depend on factors such as the type of host cells in which the recombinant polypeptide is to be produced. To illustrate, examples of heterologous signal peptides that are functional in mammalian host cells include the signal sequence for interleukin-7 (IL-7) described in United States Patent 4,965,195, the signal sequence for interleukin-2 receptor described in Cosman et al., *Nature* 312:768 (1984); the interleukin-4 receptor signal peptide described in EP 367,566; the type I interleukin-1 receptor signal peptide described in U.S. Patent 4,968,607; and the type II interleukin-1 receptor signal peptide described in EP 460,846.

Purification

The invention also includes methods of isolating and purifying the polypeptides and fragments thereof.

Isolation and Purification

The "isolated" polypeptides or fragments thereof encompassed by this invention are polypeptides or fragments that are not in an environment identical to an environment in which it or they can be found in nature. The "purified" polypeptides or fragments thereof encompassed by this invention are essentially free of association with other proteins or polypeptides, for example, as a purification product of recombinant expression systems such as those described above or as a purified product from a non-recombinant source such as naturally occurring cells and/or tissues.

In one preferred embodiment, the purification of recombinant polypeptides or fragments can be accomplished using fusions of polypeptides or fragments of the invention to another polypeptide to aid in the purification of polypeptides or fragments of the invention. Such fusion partners can include the poly-His or other antigenic identification peptides described above as well as the Fc moieties described previously.

With respect to any type of host cell, as is known to the skilled artisan, procedures for purifying a recombinant polypeptide or fragment will vary according to such factors as the type of host cells employed and whether or not the recombinant polypeptide or fragment is secreted into the culture medium.

In general, the recombinant polypeptide or fragment can be isolated from the host cells if not secreted, or from the medium or supernatant if soluble and secreted, followed by one or more concentration, salting-out, ion exchange, hydrophobic interaction, affinity purification or size exclusion chromatography steps. As to specific ways to accomplish these steps, the culture medium first can be concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. Following the concentration step, the concentrate can be applied to a purification matrix such as a gel filtration medium. Alternatively, an anion exchange resin can be employed, for example, a matrix or substrate having pendant diethylaminoethyl (DEAE) groups. The matrices can be acrylamide, agarose, dextran, cellulose or other types commonly employed in protein purification. Alternatively, a cation exchange step can be employed. Suitable cation exchangers include various insoluble matrices comprising sulfopropyl or carboxymethyl groups. In addition, a chromatofocusing step can be employed. Alternatively, a hydrophobic interaction chromatography step can be employed. Suitable matrices can be phenyl or octyl moieties bound to resins. In addition, affinity chromatography with a matrix which selectively binds the recombinant protein can be

employed. Examples of such resins employed are lectin columns, dye columns, and metal-chelating columns. Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, (e.g., silica gel or polymer resin having pendant methyl, octyl, octyldecyl or other aliphatic groups) can be employed to further purify the polypeptides. Some or all of the foregoing purification steps, in various combinations, are well known and can be employed to provide an isolated and purified recombinant protein.

It is also possible to utilize an affinity column comprising a polypeptide-binding protein of the invention, such as a monoclonal antibody generated against polypeptides of the invention, to affinity-purify expressed polypeptides. These polypeptides can be removed from an affinity column using conventional techniques, e.g., in a high salt elution buffer and then dialyzed into a lower salt buffer for use or by changing pH or other components depending on the affinity matrix utilized, or be competitively removed using the naturally occurring substrate of the affinity moiety, such as a polypeptide derived from the invention.

In this aspect of the invention, polypeptide-binding proteins, such as the anti-polypeptide antibodies of the invention or other proteins that may interact with the polypeptide of the invention, can be bound to a solid phase support such as a column chromatography matrix or a similar substrate suitable for identifying, separating, or purifying cells that express polypeptides of the invention on their surface. Adherence of polypeptide-binding proteins of the invention to a solid phase contacting surface can be accomplished by any means. For example, magnetic microspheres can be coated with these polypeptide-binding proteins and held in the incubation vessel through a magnetic field. Suspensions of cell mixtures are contacted with the solid phase that has such polypeptide-binding proteins thereon. Cells having polypeptides of the invention on their surface bind to the fixed polypeptide-binding protein and unbound cells then are washed away. This affinity-binding method is useful for purifying, screening, or separating such polypeptide-expressing cells from solution. Methods of releasing positively selected cells from the solid phase are known in the art and encompass, for example, the use of enzymes. Such enzymes are preferably non-toxic and non-injurious to the cells and are preferably directed to cleaving the cell-surface binding partner.

Alternatively, mixtures of cells suspected of containing polypeptide-expressing cells of the invention first can be incubated with a biotinylated polypeptide-binding protein of the invention. Incubation periods are typically at least one hour in duration to ensure sufficient

binding to polypeptides of the invention. The resulting mixture then is passed through a column packed with avidin-coated beads, whereby the high affinity of biotin for avidin provides the binding of the polypeptide-binding cells to the beads. Use of avidin-coated beads is known in the art. See Berenson, et al. *J. Cell. Biochem.*, 10D:239 (1986). Wash of
5 unbound material and the release of the bound cells is performed using conventional methods.

The desired degree of purity depends on the intended use of the protein. A relatively high degree of purity is desired when the polypeptide is to be administered *in vivo*, for example. In such a case, the polypeptides are purified such that no protein bands
10 corresponding to other proteins are detectable upon analysis by SDS-polyacrylamide gel electrophoresis (SDS-PAGE). It will be recognized by one skilled in the pertinent field that multiple bands corresponding to the polypeptide may be visualized by SDS-PAGE, due to differential glycosylation, differential post-translational processing, and the like. Most preferably, the polypeptide of the invention is purified to substantial homogeneity, as
15 indicated by a single protein band upon analysis by SDS-PAGE. The protein band may be visualized by silver staining, Coomassie blue staining, or (if the protein is radiolabeled) by autoradiography.

Assays

20 The purified polypeptides of the invention (including proteins, polypeptides, fragments, variants, oligomers, and other forms) may be tested for the ability to bind a cognate, ligand, receptor, substrate, or counter-structure and the like ("binding partner") in any suitable assay, such as a conventional binding assay. To illustrate, the polypeptide may be labeled with a detectable reagent (e.g., a radionuclide, chromophore, enzyme that
25 catalyzes a colorimetric or fluorometric reaction, and the like). The labeled polypeptide is contacted with cells expressing the binding partner. The cells then are washed to remove unbound labeled polypeptide, and the presence of cell-bound label is determined by a suitable technique, chosen according to the nature of the label.

One example of a binding assay procedure is as follows. A recombinant expression
30 vector containing the binding partner cDNA is constructed using methods known in the art. CV1-EBNA-1 cells in 10 cm² dishes are transfected with the recombinant expression vector. CV1-EBNA-1 cells (ATCC CRL 10478) constitutively express EBV nuclear antigen-1 driven from the CMV immediate-early enhancer/promoter. CV1-EBNA-1 was derived from

the African Green Monkey kidney cell line CV-1 (ATCC CCL 70) described by McMahan et al. (*EMBO J.* 10:2821, 1991).

The transfected cells are cultured for 24 hours, and the cells in each dish then are split into a 24-well plate. After culturing an additional 48 hours, the transfected cells (about
5 4×10^4 cells well) are washed with BM-NFDM, which is binding medium (RPMI 1640 containing 25 mg/ml bovine serum albumin, 2 mg/ml sodium azide, 20 mM Hepes pH 7.2) to which 50 mg/ml nonfat dry milk has been added. The cells then are incubated for 1 hour at 37°C with various concentrations of, for example, a soluble polypeptide/Fc fusion protein made as set forth above. Cells then are washed and incubated with a constant saturating
10 concentration of a ^{125}I -mouse anti-human IgG in binding medium, with gentle agitation for 1 hour at 37°C. After extensive washing, cells are released *via* trypsinization.

The mouse anti-human IgG employed above is directed against the Fc region of human IgG and can be obtained from Jackson ImmunoResearch Laboratories, Inc., West Grove, PA. The antibody is radioiodinated using the standard chloramine-T method. The
15 antibody will bind to the Fc portion of any polypeptide/Fc protein that has bound to the cells. In all assays, non-specific binding of ^{125}I -antibody is assayed in the absence of the Fc fusion protein/Fc, as well as in the presence of the Fc fusion protein and a 200-fold molar excess of unlabeled mouse anti-human IgG antibody.

Cell-bound ^{125}I -antibody is quantified on a Packard Autogamma counter. Affinity
20 calculations (Scatchard, *Ann. N.Y. Acad. Sci.* 51:660, 1949) are generated on RS/1 (BBN Software, Boston, MA) run on a Microvax computer.

Another type of suitable binding assay is a competitive binding assay. To illustrate, biological activity of a variant may be determined by assaying for the variant's ability to compete with the native protein for binding to the binding partner.

Competitive binding assays can be performed by conventional methodology.
25 Reagents that may be employed in competitive binding assays include radiolabeled IMX polypeptides and intact cells expressing the IMX polypeptide (endogenous or recombinant) on the cell surface. For example, a radiolabeled soluble IMX polypeptide fragment can be used to compete with a soluble IMX polypeptide variant for binding to the cell surface
30 binding partner. Instead of intact cells, one could substitute a soluble binding partner/Fc fusion protein bound to a solid phase through the interaction of Protein A or Protein G (on the solid phase) with the Fc moiety. Chromatography columns that contain Protein A and Protein G include those available from Pharmacia Biotech, Inc., Piscataway, NJ.

Another type of competitive binding assay utilizes the radiolabeled soluble binding partner, such as a soluble binding partner-Fc fusion protein, and intact cells expressing the IMX polypeptide. Qualitative results can be obtained by competitive autoradiographic plate binding assays, while Scatchard plots (Scatchard, *Ann. N.Y. Acad. Sci.* 51:660, 1949) may
5 be utilized to generate quantitative results.

USE OF IMX NUCLEIC ACIDS OR OLIGONUCLEOTIDES

In addition to being used to express polypeptides as described above, the nucleic acids of the invention, including DNA, and oligonucleotides thereof can be used:

- 10 ⊃ as probes to identify nucleic acid encoding proteins homologous to IMX polypeptides;
- ⊃ to identify human chromosomes;
- ⊃ to map genes on human chromosome numbers 7, 19, and 22;
- ⊃ to identify genes associated with certain diseases, syndromes, or other
15 conditions associated with human chromosome numbers 7, 19, and 22;
- ⊃ as single-stranded sense or antisense oligonucleotides, to inhibit expression of polypeptide encoded by the IMX sequences;
- ⊃ to help detect defective genes in an individual; and
- 20 ⊃ for gene therapy.

Probes

Among the uses of nucleic acids of the invention is the use of fragments as probes or primers. Such fragments generally comprise at least about 17 contiguous nucleotides of a
25 DNA sequence. In other embodiments, a DNA fragment comprises at least 30, or at least 60, contiguous nucleotides of a DNA sequence.

Because homologs of SEQ ID NOs:1-23, from other mammalian species, are contemplated herein, probes based on the human DNA sequence of SEQ ID NOs:1-23 may be used to screen cDNA libraries derived from other mammalian species, using
30 conventional cross-species hybridization techniques.

Using knowledge of the genetic code in combination with the amino acid sequences set forth above, sets of degenerate oligonucleotides can be prepared. Such oligonucleotides

are useful as primers, e.g., in polymerase chain reactions (PCR), whereby DNA fragments are isolated and amplified.

Identifying Chromosome Number

5 All or a portion of the nucleic acids of SEQ ID NOs:1-23, including oligonucleotides, can be used by those skilled in the art using well-known techniques to identify the human chromosomes, and the specific locus thereof, that contain the DNA of IMX family members. Useful techniques include, but are not limited to, using the sequence or portions, including oligonucleotides, as a probe in various well-known techniques such as
10 in situ hybridization to chromosome spreads, Southern blot hybridization to hybrid cell lines, fluorescent tagging, and radiation hybrid mapping.

For example, chromosomes can be mapped by radiation hybridization. PCR is performed using the Whitehead Institute/MIT Center for Genome Research Genebridge4 panel of 93 radiation hybrids ([http://www-genome.wi.mit.edu/ftp/distribution/](http://www-genome.wi.mit.edu/ftp/distribution/human_STS_releases/july97/rhmap/genebridge4.html)
15 [human_STS_releases/july97/rhmap/genebridge4.html](http://www-genome.wi.mit.edu/ftp/distribution/human_STS_releases/july97/rhmap/genebridge4.html)). Primers are used which lie within a putative exon of the gene of interest and which amplify a product from human genomic DNA, but do not amplify hamster genomic DNA. The results of the PCRs are converted into a data vector that is submitted to the Whitehead/MIT Radiation Mapping site on the internet (<http://www-seq.wi.mit.edu>). The data is scored and the chromosomal assignment
20 and placement relative to known Sequence Tag Site (STS) markers on the radiation hybrid map is provided. The following web site provides additional information about radiation hybrid mapping:

http://www-genome.wi.mit.edu/ftp/distribution/human_STS_releases/july97/07-97.INTRO.html).

25

Identifying Associated Diseases

As described previously, IMX molecules numbered 4, 21, 44, and 56 have been mapped to particular chromosome locations. Thus, the nucleic acid of a particular IMX molecule or a fragment thereof can be used by one skilled in the art using well-known
30 techniques to analyze abnormalities associated with gene mapping to such chromosomes. This enables one to distinguish conditions in which this marker is rearranged or deleted. In addition, nucleotides of such IMX molecules or fragments thereof can be used as a positional marker to map other genes of previously unknown location.

The DNA may be used in developing treatments for any disorder mediated (directly or indirectly) by defective, or insufficient amounts of, the genes corresponding to the nucleic acids of the invention. Disclosure herein of native nucleotide sequences permits the detection of defective genes, and the replacement thereof with normal genes. Defective genes may be detected in *in vitro* diagnostic assays, and by comparison of a native nucleotide sequence disclosed herein with that of a gene derived from a person suspected of harboring a defect in this gene.

Sense-Antisense

Other useful fragments of the nucleic acids include antisense or sense oligonucleotides comprising a single-stranded nucleic acid sequence (either RNA or DNA) capable of binding to target mRNA (sense) or DNA (antisense) sequences. Antisense or sense oligonucleotides according to the present invention comprise a fragment of DNA (SEQ ID NOs:1-23). Such a fragment generally comprises at least about 14 nucleotides, preferably from about 14 to about 30 nucleotides. The ability to derive an antisense or a sense oligonucleotide, based upon a cDNA sequence encoding a given protein is described in, for example, Stein and Cohen (*Cancer Res.* 48:2659, 1988) and van der Krol et al. (*BioTechniques* 6:958, 1988).

Binding of antisense or sense oligonucleotides to target nucleic acid sequences results in the formation of duplexes that block or inhibit protein expression by one of several means, including enhanced degradation of the mRNA by RNaseH, inhibition of splicing, premature termination of transcription or translation, or by other means. The antisense oligonucleotides thus may be used to block expression of proteins. Antisense or sense oligonucleotides further comprise oligonucleotides having modified sugar-phosphodiester backbones (or other sugar linkages, such as those described in WO91/06629) and wherein such sugar linkages are resistant to endogenous nucleases. Such oligonucleotides with resistant sugar linkages are stable *in vivo* (*i.e.*, capable of resisting enzymatic degradation) but retain sequence specificity to be able to bind to target nucleotide sequences.

Other examples of sense or antisense oligonucleotides include those oligonucleotides which are covalently linked to organic moieties, such as those described in WO 90/10448, and other moieties that increases affinity of the oligonucleotide for a target nucleic acid sequence, such as poly-(L-lysine). Further still, intercalating agents, such as

ellipticine, and alkylating agents or metal complexes may be attached to sense or antisense oligonucleotides to modify binding specificities of the antisense or sense oligonucleotide for the target nucleotide sequence.

Antisense or sense oligonucleotides may be introduced into a cell containing the target nucleic acid sequence by any gene transfer method, including, for example, lipofection, CaPO₄-mediated DNA transfection, electroporation, or by using gene transfer vectors such as Epstein-Barr virus.

Sense or antisense oligonucleotides also may be introduced into a cell containing the target nucleotide sequence by formation of a conjugate with a ligand binding molecule, as described in WO 91/04753. Suitable ligand binding molecules include, but are not limited to, cell surface receptors, growth factors, other cytokines, or other ligands that bind to cell surface receptors. Preferably, conjugation of the ligand binding molecule does not substantially interfere with the ability of the ligand binding molecule to bind to its corresponding molecule or receptor, or block entry of the sense or antisense oligonucleotide or its conjugated version into the cell.

Alternatively, a sense or an antisense oligonucleotide may be introduced into a cell containing the target nucleic acid sequence by formation of an oligonucleotide-lipid complex, as described in WO 90/10448. The sense or antisense oligonucleotide-lipid complex is preferably dissociated within the cell by an endogenous lipase.

USE OF IMX POLYPEPTIDES AND FRAGMENTED POLYPEPTIDES

Uses include, but are not limited to, the following:

- Purifying proteins and measuring activity thereof
- Delivery Agents
- Therapeutic and Research Reagents
- Molecular weight and Isoelectric focusing markers
- Controls for peptide fragmentation
- Identification of unknown proteins
- Preparation of Antibodies

Purification Reagents

Each of the polypeptides of the invention finds use as a protein purification reagent. The polypeptides may be attached to a solid support material and used to purify (binding partner) proteins by affinity chromatography. In particular embodiments, a polypeptide (in any form described herein that is capable of binding (binding partner)) is attached to a solid support by conventional procedures. As one example, chromatography columns containing functional groups that will react with functional groups on amino acid side chains of proteins are available (Pharmacia Biotech, Inc., Piscataway, NJ). In an alternative, a polypeptide/Fc protein (as discussed above) is attached to Protein A- or Protein G-containing chromatography columns through interaction with the Fc moiety.

The polypeptide also finds use in purifying or identifying cells that express (binding partner) on the cell surface. Polypeptides are bound to a solid phase such as a column chromatography matrix or a similar suitable substrate. For example, magnetic microspheres can be coated with the polypeptides and held in an incubation vessel through a magnetic field. Suspensions of cell mixtures containing (binding partner) expressing cells are contacted with the solid phase having the polypeptides thereon. Cells expressing (binding partner) on the cell surface bind to the fixed polypeptides, and unbound cells then are washed away.

Alternatively, the polypeptides can be conjugated to a detectable moiety, then incubated with cells to be tested for (binding partner) expression. After incubation, unbound labeled matter is removed and the presence or absence of the detectable moiety on the cells is determined.

In a further alternative, mixtures of cells suspected of containing (binding partner) cells are incubated with biotinylated polypeptides. Incubation periods are typically at least one hour in duration to ensure sufficient binding. The resulting mixture then is passed through a column packed with avidin-coated beads, whereby the high affinity of biotin for avidin provides binding of the desired cells to the beads. Procedures for using avidin-coated beads are known (see Berenson, et al. *J. Cell. Biochem.*, 10D:239, 1986). Washing to remove unbound material, and the release of the bound cells, are performed using conventional methods.

Measuring Activity

Polypeptides also find use in measuring the biological activity of (binding partner) protein in terms of their binding affinity. The polypeptides thus may be employed by those conducting "quality assurance" studies, e.g., to monitor shelf life and stability of protein under different conditions. For example, the polypeptides may be employed in a binding affinity study to measure the biological activity of a (binding partner) protein that has been stored at different temperatures, or produced in different cell types. The proteins also may be used to determine whether biological activity is retained after modification of a (binding partner) protein (e.g., chemical modification, truncation, mutation, etc.). The binding affinity of the modified (binding partner) protein is compared to that of an unmodified (binding partner) protein to detect any adverse impact of the modifications on biological activity of (binding partner). The biological activity of a (binding partner) protein thus can be ascertained before it is used in a research study, for example.

Delivery Agents

The polypeptides also find use as carriers for delivering agents attached thereto to cells bearing binding partner. Cells expressing (binding partner) include those identified in (add citation if reference known). The polypeptides thus can be used to deliver diagnostic or therapeutic agents to such cells (or to other cell types found to express (binding partner) on the cell surface) in *in vitro* or *in vivo* procedures.

Detectable (diagnostic) and therapeutic agents that may be attached to a polypeptide include, but are not limited to, toxins, other cytotoxic agents, drugs, radionuclides, chromophores, enzymes that catalyze a colorimetric or fluorometric reaction, and the like, with the particular agent being chosen according to the intended application. Among the toxins are ricin, abrin, diphtheria toxin, *Pseudomonas aeruginosa* exotoxin A, ribosomal inactivating proteins, mycotoxins such as trichothecenes, and derivatives and fragments (e.g., single chains) thereof. Radionuclides suitable for diagnostic use include, but are not limited to, ^{123}I , ^{131}I , $^{99\text{m}}\text{Tc}$, ^{111}In , and ^{76}Br . Examples of radionuclides suitable for therapeutic use are ^{131}I , ^{211}At , ^{77}Br , ^{186}Re , ^{188}Re , ^{212}Pb , ^{212}Bi , ^{109}Pd , ^{64}Cu , and ^{67}Cu .

Such agents may be attached to the polypeptide by any suitable conventional procedure. The polypeptide comprises functional groups on amino acid side chains that can be reacted with functional groups on a desired agent to form covalent bonds, for example.

Alternatively, the protein or agent may be derivatized to generate or attach a desired reactive functional group. The derivatization may involve attachment of one of the bifunctional coupling reagents available for attaching various molecules to proteins (Pierce Chemical Company, Rockford, Illinois). A number of techniques for radiolabeling proteins are known. Radionuclide metals may be attached to polypeptides by using a suitable bifunctional chelating agent, for example.

Conjugates comprising polypeptides and a suitable diagnostic or therapeutic agent (preferably covalently linked) are thus prepared. The conjugates are administered or otherwise employed in an amount appropriate for the particular application.

Therapeutic Agents

Polypeptides of the invention may be used in developing treatments for any disorder mediated (directly or indirectly) by defective, or insufficient amounts of the polypeptides. These polypeptides may be administered to a mammal afflicted with such a disorder.

The polypeptides may also be employed in inhibiting a biological activity of (binding partner), in *in vitro* or *in vivo* procedures. For example, a purified polypeptide may be used to inhibit binding of (binding partner) to endogenous cell surface (binding partner). Biological effects that result from the binding of (binding partner) to endogenous receptors thus are inhibited.

IMX polypeptides may be administered to a mammal to treat a (binding partner)-mediated disorder. Such (binding partner)-mediated disorders include conditions caused (directly or indirectly) or exacerbated by (binding partner).

Compositions of the present invention may contain a polypeptide in any form described herein, such as native proteins, variants, derivatives, oligomers, and biologically active fragments. In particular embodiments, the composition comprises a soluble polypeptide or an oligomer comprising soluble polypeptides.

Compositions comprising an effective amount of a polypeptide of the present invention, in combination with other components such as a physiologically acceptable diluent, carrier, or excipient, are provided herein. The polypeptides can be formulated according to known methods used to prepare pharmaceutically useful compositions. They can be combined in admixture, either as the sole active material or with other known active materials suitable for a given indication, with pharmaceutically acceptable diluents (e.g., saline, Tris-HCl, acetate, and phosphate buffered solutions), preservatives (e.g., thimerosal,

benzyl alcohol, parabens), emulsifiers, solubilizers, adjuvants and or carriers. Suitable formulations for pharmaceutical compositions include those described in *Remington's Pharmaceutical Sciences*, 16th ed. 1980, Mack Publishing Company, Easton, PA.

5 In addition, such compositions can be complexed with polyethylene glycol (PEG), metal ions, or incorporated into polymeric compounds such as polyacetic acid, polyglycolic acid, hydrogels, dextran, etc., or incorporated into liposomes, microemulsions, micelles, unilamellar or multilamellar vesicles, erythrocyte ghosts or spheroblasts. Such compositions will influence the physical state, solubility, stability, rate of *in vivo* release, and rate of *in vivo* clearance, and are thus chosen according to the intended application.

10 The compositions of the invention can be administered in any suitable manner, e.g., topically, parenterally, or by inhalation. The term "parenteral" includes injection, e.g., by subcutaneous, intravenous, or intramuscular routes, also including localized administration, e.g., at a site of disease or injury. Sustained release from implants is also contemplated. One skilled in the pertinent art will recognize that suitable dosages will vary, depending
15 upon such factors as the nature of the disorder to be treated, the patient's body weight, age, and general condition, and the route of administration. Preliminary doses can be determined according to animal tests, and the scaling of dosages for human administration is performed according to art-accepted practices.

20 Compositions comprising nucleic acids in physiologically acceptable formulations are also contemplated. DNA may be formulated for injection, for example.

Research Agents

Another use of the polypeptide of the present invention is as a research tool for studying the biological effects that result from inhibiting interactions between polypeptides
25 of the invention and their "binding partner". Polypeptides also may be employed in *in vitro* assays for detecting binding partner or cells expressing the binding partner or the interactions thereof.

Molecular Weight, Isoelectric Point Markers

The polypeptides of the present invention can be subjected to fragmentation into smaller peptides by chemical and enzymatic means, and the peptide fragments so produced can be used in the analysis of other proteins or polypeptides. For example, such peptide
5 fragments can be used as peptide molecular weight markers, peptide isoelectric point markers, or in the analysis of the degree of peptide fragmentation. Thus, the invention also includes these polypeptides and peptide fragments, as well as kits to aid in the determination of the apparent molecular weight and isoelectric point of an unknown protein and kits to assess the degree of fragmentation of an unknown protein.

10 Although all methods of fragmentation are encompassed by the invention, chemical fragmentation is a preferred embodiment, and includes the use of cyanogen bromide to cleave under neutral or acidic conditions such that specific cleavage occurs at methionine residues (E. Gross, *Methods in Enz.* 11:238-255, 1967). This can further include additional steps, such as a carboxymethylation step to convert cysteine residues to an unreactive
15 species.

Enzymatic fragmentation is another preferred embodiment, and includes the use of a protease such as Asparaginylendo-peptidase, Arginylendo-peptidase, *Achromobacter* protease I, Trypsin, *Staphylococcus aureus* V8 protease, Endoproteinase Asp-N, or Endoproteinase Lys-C under conventional conditions to result in cleavage at specific amino
20 acid residues. Asparaginylendo-peptidase can cleave specifically on the carboxyl side of the asparagine residues present within the polypeptides of the invention. Arginylendo-peptidase can cleave specifically on the carboxyl side of the arginine residues present within these polypeptides. *Achromobacter* protease I can cleave specifically on the carboxyl side of the lysine residues present within the polypeptides (Sakiyama and Nakat, U.S. Patent
25 No. 5,248,599; T. Masaki et al., *Biochim. Biophys. Acta* 660:44-50, 1981; T. Masaki et al., *Biochim. Biophys. Acta* 660:51-55, 1981). Trypsin can cleave specifically on the carboxyl side of the arginine and lysine residues present within polypeptides of the invention. Enzymatic fragmentation may also occur with a protease that cleaves at multiple amino acid residues. For example, *Staphylococcus aureus* V8 protease can cleave specifically on the
30 carboxyl side of the aspartic and glutamic acid residues present within polypeptides (D. W. Cleveland, *J. Biol. Chem.* 3:1102-1106, 1977). Endoproteinase Asp-N can cleave specifically on the amino side of the asparagine residues present within polypeptides. Endoproteinase Lys-C can cleave specifically on the carboxyl side of the lysine residues

present within polypeptides of the invention. Other enzymatic and chemical treatments can likewise be used to specifically fragment these polypeptides into a unique set of specific peptides.

Of course, the peptides and fragments of the polypeptides of the invention can also be produced by conventional recombinant processes and synthetic processes well known in the art. With regard to recombinant processes, the polypeptides and peptide fragments encompassed by invention can have variable molecular weights, depending upon the host cell in which they are expressed. Glycosylation of polypeptides and peptide fragments of the invention in various cell types can result in variations of the molecular weight of these pieces, depending upon the extent of modification. The size of these pieces can be most heterogeneous with fragments of polypeptide derived from the extracellular portion of the polypeptide. Consistent polypeptides and peptide fragments can be obtained by using polypeptides derived entirely from the transmembrane and cytoplasmic regions, pretreating with N-glycanase to remove glycosylation, or expressing the polypeptides in bacterial hosts.

The molecular weight of these polypeptides can also be varied by fusing additional peptide sequences to both the amino and carboxyl terminal ends of polypeptides of the invention. Fusions of additional peptide sequences at the amino and carboxyl terminal ends of polypeptides of the invention can be used to enhance expression of these polypeptides or aid in the purification of the protein. In addition, fusions of additional peptide sequences at the amino and carboxyl terminal ends of polypeptides of the invention will alter some, but usually not all, of the fragmented peptides of the polypeptides generated by enzymatic or chemical treatment. Of course, mutations can be introduced into polypeptides of the invention using routine and known techniques of molecular biology. For example, a mutation can be designed so as to eliminate a site of proteolytic cleavage by a specific enzyme or a site of cleavage by a specific chemically induced fragmentation procedure. The elimination of the site will alter the peptide fingerprint of polypeptides of the invention upon fragmentation with the specific enzyme or chemical procedure.

The polypeptides and the resultant fragmented peptides can be analyzed by methods including sedimentation, electrophoresis, chromatography, and mass spectrometry to determine their molecular weights. Because the unique amino acid sequence of each piece specifies a molecular weight, these pieces can thereafter serve as molecular weight markers using such analysis techniques to assist in the determination of the molecular weight of an unknown protein, polypeptides or fragments thereof. The molecular weight markers of the

invention serve particularly well as molecular weight markers for the estimation of the apparent molecular weight of proteins that have similar apparent molecular weights and, consequently, allow increased accuracy in the determination of apparent molecular weight of proteins.

5 When the invention relates to the use of fragmented peptide molecular weight markers, those markers are preferably at least 10 amino acids in size. More preferably, these fragmented peptide molecular weight markers are between 10 and 100 amino acids in size. Even more preferable are fragmented peptide molecular weight markers between 10 and 50 amino acids in size and especially between 10 and 35 amino acids in size. Most preferable
10 are fragmented peptide molecular weight markers between 10 and 20 amino acids in size.

 Among the methods for determining molecular weight are sedimentation, gel electrophoresis, chromatography, and mass spectrometry. A particularly preferred embodiment is denaturing polyacrylamide gel electrophoresis (U. K. Laemmli, *Nature* 227:680-685, 1970). Conventionally, the method uses two separate lanes of a gel containing
15 sodium dodecyl sulfate and a concentration of acrylamide between 6-20%. The ability to simultaneously resolve the marker and the sample under identical conditions allows for increased accuracy. It is understood, of course, that many different techniques can be used for the determination of the molecular weight of an unknown protein using polypeptides of the invention, and that this embodiment in no way limits the scope of the invention.

20 Each unglycosylated polypeptide or fragment thereof has a pI that is intrinsically determined by its unique amino acid sequence (which pI can be estimated by the skilled artisan using any of the computer programs designed to predict pI values currently available, calculated using any well-known amino acid pKa table, or measured empirically). Therefore these polypeptides and fragments thereof can serve as specific markers to assist in
25 the determination of the isoelectric point of an unknown protein, polypeptide, or fragmented peptide using techniques such as isoelectric focusing. These polypeptide or fragmented peptide markers serve particularly well for the estimation of apparent isoelectric points of unknown proteins that have apparent isoelectric points close to that of the polypeptide or fragmented peptide markers of the invention.

30 The technique of isoelectric focusing can be further combined with other techniques such as gel electrophoresis to simultaneously separate a protein on the basis of molecular weight and charge. The ability to simultaneously resolve these polypeptide or fragmented peptide markers and the unknown protein under identical conditions allows for increased

accuracy in the determination of the apparent isoelectric point of the unknown protein. This is of particular interest in techniques, such as two dimensional electrophoresis (T.D. Brock and M.T. Madigan, *Biology of Microorganisms* 76-77 (Prentice Hall, 6d ed. 1991)), where the nature of the procedure dictates that any markers should be resolved simultaneously with the unknown protein. In addition, with such methods, these polypeptides and fragmented peptides thereof can assist in the determination of both the isoelectric point and molecular weight of an unknown protein or fragmented peptide.

Polypeptides and fragmented peptides can be visualized using two different methods that allow a discrimination between the unknown protein and the molecular weight markers.

In one embodiment, the polypeptide and fragmented peptide molecular weight markers of the invention can be visualized using antibodies generated against these markers and conventional immunoblotting techniques. This detection is performed under conventional conditions that do not result in the detection of the unknown protein. It is understood that it may not be possible to generate antibodies against all polypeptide fragments of the invention, since small peptides may not contain immunogenic epitopes. It is further understood that not all antibodies will work in this assay; however, those antibodies which are able to bind polypeptides and fragments of the invention can be readily determined using conventional techniques.

The unknown protein is also visualized by using a conventional staining procedure.

The molar excess of unknown protein to polypeptide or fragmented peptide molecular weight markers of the invention is such that the conventional staining procedure predominantly detects the unknown protein. The level of these polypeptide or fragmented peptide molecular weight markers is such as to allow little or no detection of these markers by the conventional staining method. The preferred molar excess of unknown protein to polypeptide molecular weight markers of the invention is between 2 and 100,000 fold. More preferably, the preferred molar excess of unknown protein to these polypeptide molecular weight markers is between 10 and 10,000 fold and especially between 100 and 1,000 fold.

It is understood of course that many techniques can be used for the determination and detection of molecular weight and isoelectric point of an unknown protein, polypeptides, and fragmented peptides thereof using these polypeptide molecular weight markers and peptide fragments thereof and that these embodiments in no way limit the scope of the invention.

In another embodiment, the analysis of the progressive fragmentation of the polypeptides of the invention into specific peptides (D. W. Cleveland et al., *J. Biol. Chem.* 252:1102-1106, 1977), such as by altering the time or temperature of the fragmentation reaction, can be used as a control for the extent of cleavage of an unknown protein. For example, cleavage of the same amount of polypeptide and unknown protein under identical conditions can allow for a direct comparison of the extent of fragmentation. Conditions that result in the complete fragmentation of the polypeptide can also result in complete fragmentation of the unknown protein.

As to the specific use of the polypeptides and fragmented peptides of the invention as molecular weight markers, the fragmentation of the polypeptides of SEQ ID NOs:24-33 with cyanogen bromide generates a unique set of fragmented peptide molecular weight markers. An additional fragment results if the initiating methionine is present. The distribution of methionine residues determines the number of amino acids in each peptide and the unique amino acid composition of each peptide determines its molecular weight.

In addition, the preferred purified polypeptides of the invention (SEQ ID NOs:24-33) have calculated molecular weights of approximately 3683, 1783, 11248, 75503, 43040, 8051, 33306, 3515, 10736, 25162, and 2450 Daltons.

Where an intact protein is used, the use of these polypeptide molecular weight markers allows increased accuracy in the determination of apparent molecular weight of proteins that have apparent molecular weights close to 3683, 1783, 11248, 75503, 43040, 8051, 33306, 3515, 10736, 25162, or 2450 Daltons. Where fragments are used, there is increased accuracy in determining molecular weight over the range of the molecular weights of the fragment.

Finally, as to the kits that are encompassed by the invention, the constituents of such kits can be varied, but typically contain the polypeptide and fragmented peptide molecular weight markers. Also, such kits can contain the polypeptides wherein a site necessary for fragmentation has been removed. Furthermore, the kits can contain reagents for the specific cleavage of the polypeptide and the unknown protein by chemical or enzymatic cleavage. Kits can further contain antibodies directed against polypeptides or fragments thereof of the invention.

Identification of Unknown Proteins

As set forth above, a polypeptide or peptide fingerprint can be entered into or compared to a database of known proteins to assist in the identification of the unknown protein using mass spectrometry (W.J. Henzel et al., *Proc. Natl. Acad. Sci. USA* 90:5011-5015, 1993; D. Fenyo et al., *Electrophoresis* 19:99S-100S, 1998). A variety of computer software programs to facilitate these comparisons are accessible via the Internet, such as Protein Prospector (Internet site: prospector.uscf.edu), MultiIdent (Internet site: www.expasy.ch/spot/multiident.html), PeptideSearch (Internet site: www.mann.embl-heidelberg.de/deSearch/FR_PeptideSearch_Form.html), and ProFound (Internet site: www.chait-sgi.rockefeller.edu/cgi-bin/prot-id-frag.html). These programs allow the user to specify the cleavage agent and the molecular weights of the fragmented peptides within a designated tolerance. The programs compare these molecular weights to protein databases to assist in determining the identity of the unknown protein.

In addition, a polypeptide or peptide digest can be sequenced using tandem mass spectrometry (MS/MS) and the resulting sequence searched against databases (J.K. Eng, et al., *J. Am. Soc. Mass Spec.* 5:976-989 (1994); M. Mann and M. Wilm, *Anal. Chem.* 66:4390-4399 (1994); J.A. Taylor and R.S. Johnson, *Rapid Comm. Mass Spec.* 11:1067-1075 (1997)). Searching programs that can be used in this process exist on the Internet, such as Lutefisk 97 (Internet site: www.isbc.com:70/Lutefisk97.html), and the Protein Prospector, Peptide Search and ProFound programs described above. Therefore, adding the sequence of a gene and its predicted protein sequence and peptide fragments to a sequence database can aid in the identification of unknown proteins using tandem mass spectrometry.

Antibodies

Antibodies that are immunoreactive with the polypeptides of the invention are provided herein. Such antibodies specifically bind to the polypeptides *via* the antigen-binding sites of the antibody (as opposed to non-specific binding). Thus, the polypeptides, fragments, variants, fusion proteins, etc., as set forth above may be employed as immunogens in producing antibodies immunoreactive therewith.

Polyclonal and monoclonal antibodies may be prepared by conventional techniques. See, for example, *Monoclonal Antibodies, Hybridomas: A New Dimension in Biological Analyses*, Kennet et al. (eds.), Plenum Press, New York (1980); and *Antibodies: A Laboratory Manual*, Harlow and Land (eds.), Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, (1988).

Antigen-binding fragments of such antibodies, which may be produced by conventional techniques, are also encompassed by the present invention. Examples of such fragments include, but are not limited to, Fab and F(ab')₂ fragments. Antibody fragments and derivatives produced by genetic engineering techniques are also provided.

5 The monoclonal antibodies of the present invention include chimeric antibodies, e.g., humanized versions of murine monoclonal antibodies. Such humanized antibodies may be prepared by known techniques, and offer the advantage of reduced immunogenicity when the antibodies are administered to humans. In one embodiment, a humanized monoclonal antibody comprises the variable region of a murine antibody (or just the antigen
10 binding site thereof) and a constant region derived from a human antibody. Alternatively, a humanized antibody fragment may comprise the antigen binding site of a murine monoclonal antibody and a variable region fragment (lacking the antigen-binding site) derived from a human antibody. Procedures for the production of chimeric and further engineered monoclonal antibodies include those described in Riechmann et al. (*Nature*
15 332:323, 1988), Liu et al. (*PNAS* 84:3439, 1987), Larrick et al. (*Bio/Technology* 7:934, 1989), and Winter and Harris (*TIPS* 14:139, May, 1993). Procedures to generate antibodies transgenically can be found in GB 2,272,440, US Patent Nos. 5,569,825 and 5,545,806 and related patents claiming priority therefrom, all of which are incorporated by reference herein.

20 In one embodiment, the antibodies are specific for the polypeptides of the present invention, and do not cross-react with other proteins. Screening procedures by which such antibodies may be identified are well known, and may involve immunoaffinity chromatography, for example.

Hybridoma cell lines that produce monoclonal antibodies specific for the
25 polypeptides of the invention are also contemplated herein. Such hybridomas may be produced and identified by conventional techniques. One method for producing such a hybridoma cell line comprises immunizing an animal with a polypeptide; harvesting spleen cells from the immunized animal; fusing said spleen cells to a myeloma cell line, thereby generating hybridoma cells; and identifying a hybridoma cell line that produces a
30 monoclonal antibody that binds the polypeptide. The monoclonal antibodies may be recovered by conventional techniques.

Uses Thereof

The antibodies of the invention can be used in assays to detect the presence of the polypeptides or fragments of the invention, either *in vitro* or *in vivo*. The antibodies also may be employed in purifying polypeptides or fragments of the invention by immunoaffinity chromatography.

5 Those antibodies that additionally can block binding of the polypeptides of the invention to the binding partner may be used to inhibit a biological activity that results from such binding. Such blocking antibodies may be identified using any suitable assay procedure, such as by testing antibodies for the ability to inhibit binding of the binding partner to certain cells expressing the binding partner. Alternatively, blocking antibodies
10 may be identified in assays for the ability to inhibit a biological effect that results from binding of the binding partner to target cells.

Such an antibody may be employed in an *in vitro* procedure, or administered *in vivo* to inhibit a biological activity mediated by the entity that generated the antibody. Disorders caused or exacerbated (directly or indirectly) by the interaction of (binding partner) with cell
15 surface (binding partner) receptor thus may be treated. A therapeutic method involves *in vivo* administration of a blocking antibody to a mammal in an amount effective in inhibiting a (binding partner)-mediated biological activity. Monoclonal antibodies are generally preferred for use in such therapeutic methods. In one embodiment, an antigen-binding antibody fragment is employed.

20 Antibodies may be screened for agonistic (*i.e.*, ligand-mimicking) properties. Such antibodies, upon binding to cell surface antigen, induce biological effects (e.g., transduction of biological signals) similar to the biological effects induced when (binding partner) binds to cell surface antigen.

Compositions comprising an antibody that is directed against a polypeptide of the invention, and a physiologically acceptable diluent, excipient, or carrier, are provided
25 herein.

Also provided herein are conjugates comprising a detectable (e.g., diagnostic) or therapeutic agent, attached to the antibody. Examples of such agents are presented above. The conjugates find use in *in vitro* or *in vivo* procedures.

30 The following examples are provided to further illustrate particular embodiments of the invention, and are not to be construed as limiting the scope of the present invention.

EXAMPLE 1: Isolation of the IMX Nucleic Acids

The T84 Epithelial Barrier Model

As discussed above, damage to the intestinal epithelial barrier is a hallmark of (IBD), and a number of *in vitro* models of epithelial barrier function have been developed over the years. The best characterized of these models is the T84 intestinal epithelial barrier system, Dharmasathaporn et al., *Am. J. Physiol.*, 246:G204-G208, 1984 and Madara et al., *J. Cell Biol.*, 101:2124-2133, 1985).

T84 cells were plated on 75 mm polycarbonate transwell filter inserts (Costar) and grown in DME/F12 (1:1) containing 10% heat-inactivated bovine calf serum. The cells were maintained at confluence for 2-3 days, and integrity of the epithelial barrier was determined by measuring transepithelial electrical resistance (TER) using an EVOM epithelial voltohmmeter (World Precision Instruments). When the TER values were greater than 1000 ohms/cm² and were stable, cells were treated with interferon- γ (30 ng/ml, Genzyme) added to the basolateral side of the membrane. At various times after treatment (4, 24 and 44h), TERs were measured to monitor the interferon-induced disruption of the barrier, and RNA was harvested from the cells at those time points using TRIzol reagent (Life Technologies). RNA was extracted using conventional methods and subsequently used for TOGATM analysis as described in Example 2.

TOGATM analysis

This example describes a method for determining mRNA expression characteristics. The isolated RNA was analyzed using a method of simultaneous sequence-specific identification of mRNAs known as TOGATM (Total Gene expression Analysis) described in U.S. Patent No. 5,459,037 and U.S. Patent No. 5,807,680, hereby incorporated herein by reference. Preferably, prior to the application of the TOGATM technique, the isolated RNA was enriched to form a starting polyA-containing mRNA population by methods known in the art. In a preferred embodiment, the TOGATM method further comprised an additional PCR step performed using four separate reactions, one for each of the four 5' PCR primers, and cDNA templates prepared from a population of antisense cRNAs. A final PCR step used 256 5' PCR primers in 64 subpools for each of the four reactions of the previous step produced PCR products that were cDNA fragments that corresponded to the 3'-region of the starting mRNA population.

The produced PCR products were then identified by a) the sequence of at least the 5' seven base pairs, preferably the sequence of the entire fragment, and b) the length of the fragment. These two parameters, sequence and fragment length, were used to compare the

obtained PCR products to a database of known polynucleotide sequences. A database search for homologous sequences in Genbank resulted in no matches, indicating the novelty of the MX sequences of the invention. The intensities of the PCR products were compared across the 4 input RNA samples (t=0, 4hr, 24hr, 44hr) and species that were regulated were identified and further characterized (Table 1).

The DSTs of SEQ ID NO:1-23 and fragments thereof are useful as probes to study and diagnose the changes in gene expression demonstrated by the data of Table 1.

Alternatively, polypeptides and fragments thereof that are the translations of SEQ ID NO:1-23 and fragments thereof are useful as probes to study and diagnose the changes in gene expression demonstrated by the data of Table 1.

Name	DST SEQ ID NO:	Digital Address (Msp I)	Relative PCR Fragment Amount			
			0	4hr	24hr	44hr
IMX 4	1	TAAG 187	41	94	350	299
IMX 10	14	TTCT 159	250	716	250	207
IMX 21	15	CTTC 364	35	60	330	749
IMX 28	16	CCAT 414	138	435	647	464
IMX 32	17	CCGA 374	237	539	25	32
IMX 39	19	TGGA 450	24	87	110	78
IMX 40	7	TCTA 163	38	294	67	153
IMX 42	8	GAGC 426	145	733	876	1076
IMX 44	21	CTGC 221	248	344	1499	1624
IMX 56	10	GATA 111	955	1223	2357	3450

Table 1. IMX Nucleic Acid Molecules. Relative expression levels determined by TOGA™.

5

EXAMPLE 2: Further Characterization of IMX4

Nucleic Acid Molecules and Polypeptides

An additional 569bp sequence (SEQ ID NO:11) that included DST IMX4 (SEQ ID NO:1) was obtained by anchor PCR from a T84 library. A clone approximately 2.0kb in length was then isolated from a commercial library (Origene). The sequence of the 3' end of the clone (SEQ ID NO:12) included the sequence of the IMX4 DST. The sequence of the 5' end of the clone (SEQ ID NO:13) matched part of the human ApoL gene (AF019225, Figure 21). Complete sequence of the clone was not obtained. However, comparison of 5' end of the clone suggests it represents an alternative splice product to the reported ApoL sequence, i.e., bases 1-168 match to 2 exons on PAC carrying the ApoL gene but are not included in the reported complete cDNA. See Figure 22. A polypeptide translated from IMX4 is provided in SEQ ID NO: 24.

EXAMPLE 3: Further Characterization of IMX10 and IMX21

Nucleic Acid Molecules and Polypeptides

20

Further experiments failed to show regulation of these DSTs in replications using PCR with extended primers and Northern blots. A polypeptide translated from IMX10 (SEQ ID NO:14) is provided in SEQ ID NO: 25. A polypeptide translated from IMX21 (SEQ ID NO:15) is provided in SEQ ID NO: 26.

5

EXAMPLE 4: Further Characterization of IMX28

Nucleic Acid Molecules and Polypeptides

One polypeptide translated from IMX28 is provided in SEQ ID NO: 27.

10

A GST fusion protein of IMX28 was used in studies designed to identify potential ligands for this enzyme. The GST fusion of IMX28 was prepared as follows: IMX28-8, a clone with the full length coding sequence, was excised from pGEM by Sall-NotI digestion and subcloned into Sall-NotI digested pGEX-5X-3 (Pharmacia). The pGEX-5X-3 plasmid contains the Schistosoma japonicum glutathione-S-transferase (GST) gene 5' to the Sall-
15 NotI cloning site. Insertion of the IMX28-8 DNA fragment into this cloning site results in a fusion gene containing GST-IMX28 in the proper reading frame. The complete sequence of the fusion protein consists of GST followed by a Factor Xa tripeptide cleavage site, the peptide consisting of Gly-Ile-Pro-Arg-Asn-Ser-Arg-Val-Asp-Ala-Thr that is derived partially from the pGEX vector and partially from the Sall-NotI fragment of IMX28 and
20 IMX28. The vector containing the fusion gene was electroporated into competent bacterial cells and a single colony containing the DNA, confirmed by diagnostic restriction digestion, was expanded to a 250ml culture for purification of the fusion protein.

Briefly, cells from these cultures were pelleted and resuspended in STE buffer containing a protease inhibitor cocktail and 20ug/ml hen egg lysozyme. The cells were lysed
25 by addition of 10% sarcosyl in STE (final sarcosyl concentration was 1.5%), sonicated and centrifuged to remove cell debris. The supernatant was passed over a Glutathione-Sepharose 4B column, washed sequentially with 1.5% sarcosyl in STE and then STE. The bound material, containing the GST-IMX28 fusion protein was eluted by adding 20mM Glutathione in STE. The eluted material was analyzed by SDS-PAGE to confirm the
30 presence of the GST-IMX28 fusion protein.

A recombinant adenovirus construct expressing IMX28 was used to infect T84 cells to determine effects on epithelial barrier formation. To construct adenoviral vectors with untagged full length Imx28 and NH2-terminally FLAG tagged Imx28, the pADEASY1

system described by He et al. (1998) Proc. Natl. Acad. Sci. USA 95: 2509-2514, was used. For the untagged construct a NotI fragment carrying the full coding region of Imx28 was inserted into the polylinker of the pAdtrackCMV shuttle vector. After confirmation of the correct orientation the vector was cut with the restriction enzyme PacI and co-
5 transformed into E. coli. Strain BJ5183 along with the pADEasy1 vector. Recombinants containing the Imx28 product were screened for as described and by gene specific PCR. To generate non-replicative, infectious viral particles the helper strain 293EBNA-EBNA or 293MSR was transfected with PmeI linearized recombinant vector. After 3 days virus was harvested by freeze-thaw lysis of the transfected cells. Several rounds of infection of these
10 cells with viral supernatants were performed to boost titers. The tagged version was prepared by addition of the FLAG sequence and a Gly-Gly-Gly-Gly spacer region onto the full length coding sequence of Imx28 via PCR with a 5' oligo with the incorporated nucleotides for the tag and spacer. The PCR product was cloned into a TA- vector and excised with NotI and cloned as above. Screening and viral particle production was as
15 above. No deleterious or positive effects were observed on T84 barrier function in the presence or absence of IFN-gamma.

EXAMPLE 5: Further Characterization of IMX32

Nucleic Acid Molecules and Polypeptides

20 The original clone produced by extending the IMX32 DST was 1588 bp in length (SEQ ID NO: 17). The first round of anchor PCR yielded a 670bp product that was used to produce a contig that extended the sequence 585 bases to 2173 (SEQ ID NO: 18). Another round of anchor PCR yielded the 1676 bp product, immunex32-a24.seq. which further extended the contig for Imx32 to 2834 bp. In addition, the immunex32-a24.seq product
25 revealed an alternative splice lacking bases 211-639 of IMX32-1 clone.

There is an ORF for 155 amino acids at the 5' end of the sequence (SEQ ID NO: 28 translated from SEQ ID NO: 17). BLAST analysis suggests that the ORF may contain a KRAB-like domain because it has partial matches to Zn-finger proteins which contain KRAB domains at their N-terminus such as ZNF140.

30

EXAMPLE 6: Further Characterization of IMX39

Nucleic Acid Molecules and Polypeptides

One polypeptide translated from IMX39 is provided in SEQ ID NO: 29.

IMX39 FLAG-tagged and untagged adenoviral vector versions were prepared as outlined above for Imx28 in Example 4, above. Controls for the expression were infection with other FLAG- tagged adenoviral delivered proteins such as IMX5. Based upon the predicted structure from its cDNA sequence, it was expected that the IMX39 polypeptide would be a cytoplasmic protein. However, since some secreted cytokines, such as those of the IL-1 family, lack a predicted signal sequence but are secreted from the cell, this expectation was tested experimentally in both T84 cells via adenoviral mediated transduction and via transfection into CV1EBNA cells with another expression vector, pDC412. Despite evidence for successful expression in the transfected cells, no product was found in the media as determined by FLAG western (T84 system) or 35-S radiolabeled product (CV1EBNA system). The positive control in the CV1 system was an Imx44-Fc fusion protein known to be secrete. In both systems empty vectors were used as a negative control. The attempted expression of IMX39 polypeptides had no apparent effect on barrier function in the absence or presence of IFN-gamma.

EXAMPLE 7: Further Characterization of IMX40

Nucleic Acid Molecules and Polypeptides

Anchor PCR using an Origene human small intestine library revealed a product of 1669 bp (SEQ ID NO: 20). Bases 1-235 of the DST IMX40 (SEQ ID NO:7) align with bases 1565-1669 of this sequence. The anchor PCR product has a potential 155 aa ORF from bases 185-650. A translation of SEQ ID NO:7 is given in SEQ ID NO:30. This ORF has no recognizable features, motifs, or database homologies.

EXAMPLE 8: Further Characterization of IMX42

Nucleic Acid Molecules and Polypeptides

Several attempts to screen libraries using DST IMX42 (SEQ ID NO:8) as well as anchor PCR and RACE have not produced a clone with a longer sequence. BLAST comparison against imxhutdb extended sequence approx 200 bases to 592 bases. However, the sequence extension did not increase any database matches found by comparison to the DST IMX42 sequence. The contig has a potential ORF of 134 amino acids which, while a COOH terminal extension of SEQ ID NO:31, may be a partial sequence.

EXAMPLE 9: Further Characterization of IMX44

Nucleic Acid Molecules and Polypeptides

Longer polynucleotides corresponding to DST IMX44 have been identified (SEQ ID NO:21,22). A translated polypeptide is found in SEQ ID NO:32. A soluble Fc form of IMX44 polypeptide was synthesized and used in various assays. IMX44-Fc had no effect on T84 barrier function in the absence or presence of IFN-gamma. IMX44-Fc had no effect on natural killer (NK) cell activation. IMX44-Fc had no positive hits on cognate screen assays.

A soluble FLAG polyHis form was also produced. No activity in cellular activation assays was found nor any alteration of cytokine production using this polypeptide in assays.

The expression of IMX44 in various murine models of gut inflammation was determined by Northern and array analysis. Little to no regulation of transcript was found was found in anti-CD3-induced ileitis in C57BL/6 mice, DSS-induced colitis in BALB/c mice or C57BL/6 mice, *mdr1* knock out mice with colitis, and IFN-gamma stimulated LN T cells.

EXAMPLE 10: Further Characterization of IMX56

Nucleic Acid Molecules and Polypeptides

A translated IMX56 polypeptide is found in SEQ ID NO:33. Comparison of the DST sequence IMX56 to IMAGE consortium clones extended the sequence (SEQ ID NO:23) which was 3' on sequenced PAC to the described end of human ApoL. Anchor PCR using a T84 library produced results that indicated that the IMX56 DST is derived from an alternate 3' UTR of ApoL.

EXAMPLE 11: Monoclonal Antibodies That Bind IMX Polypeptides

Monoclonal antibodies that bind the polypeptides of the invention can be prepared by methods well known in the art. Suitable immunogens that may be employed in generating such antibodies include, but are not limited to, purified IMX polypeptides or an immunogenic fragment thereof such as the extracellular domain, or fusion proteins containing IMX polypeptides (e.g., a soluble IMX 21 polypeptide/Fc fusion protein).

Purified IMX polypeptides of the invention can be used to generate monoclonal antibodies immunoreactive therewith, using conventional techniques such as those described in U.S. Patent 4,411,993. Briefly, mice are immunized with an IMX polypeptide immunogen emulsified in complete Freund's adjuvant, and injected in amounts ranging from

10-100 µg subcutaneously or intraperitoneally. Ten to twelve days later, the immunized animals are boosted with additional IMX polypeptide immunogen emulsified in incomplete Freund's adjuvant. Mice are periodically boosted thereafter on a weekly to bi-weekly immunization schedule. Serum samples are periodically taken by retro-orbital bleeding or
5 tail-tip excision to test for anti-IMX polypeptide antibodies by dot blot assay, ELISA (Enzyme-Linked Immunosorbent Assay) or inhibition of IMX polypeptide/binding partner interactions.

Following detection of an appropriate antibody titer, positive animals are provided one last intravenous injection of IMX polypeptide immunogen in saline. Three to four days
10 later, the animals are sacrificed, spleen cells harvested, and spleen cells are fused to a murine myeloma cell line, e.g., NS1 or preferably P3x63Ag8.653 (ATCC CRL 1580). Fusions generate hybridoma cells, which are plated in multiple microtiter plates in a HAT (hypoxanthine, aminopterin and thymidine) selective medium to inhibit proliferation of non-fused cells, myeloma hybrids, and spleen cell hybrids.

15 The hybridoma cells are screened by ELISA for reactivity against the purified IMX polypeptide of interest by adaptations of the techniques disclosed in Engvall et al., *Immunochem.* 8:871, 1971 and in U.S. Patent 4,703,004. A preferred screening technique is the antibody capture technique described in Beckmann et al., (*J. Immunol.* 144:4212, 1990) Positive hybridoma cells can be injected intraperitoneally into syngeneic BALB/c mice to
20 produce ascites containing high concentrations of anti-IMX polypeptide monoclonal antibodies. Alternatively, hybridoma cells can be grown *in vitro* in flasks or roller bottles by various techniques. Monoclonal antibodies produced in mouse ascites can be purified by ammonium sulfate precipitation, followed by gel exclusion chromatography. Alternatively, affinity chromatography based upon binding of antibody to Protein A or Protein G can also
25 be used, as can affinity chromatography based upon binding to the IMX polypeptide of interest.

Alternatively, IMX nucleic acid molecules or fragments thereof can be expressed to produce IMX polypeptides or fragments thereof, that can be used to make antibodies that are useful for identifying corresponding polypeptides in techniques such as western blotting,
30 immunocytochemistry, and ELISA assays using standard techniques such as those described in U.S. Patent No. 4,900,811, incorporated by reference herein.

The references cited herein are incorporated by reference herein in their entirety.

We claim:

1. An isolated nucleic acid molecule comprising a polynucleotide chosen from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:20, SEQ ID NO:21, and SEQ ID NO:23.
2. The isolated nucleic acid molecule of claim 1 comprising a polynucleotide shown in SEQ ID NO:1.
3. An isolated polypeptide chosen from the group consisting of SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:32, and SEQ ID NO:33.
4. The isolated polypeptide of claim 3 comprising a polypeptide shown in SEQ ID NO:24.
5. An isolated nucleic acid molecule comprising a polynucleotide at least 95% identical to the isolated nucleic acid molecule of claim 1.
6. An isolated nucleic acid molecule at least ten bases in length that is hybridizable to the isolated nucleic acid molecule of claim 1 under stringent conditions.
7. An isolated nucleic acid molecule encoding the polypeptide of claim 3.
8. An isolated nucleic acid molecule encoding a fragment of the polypeptide of claim 3.
9. An isolated nucleic acid molecule encoding a polypeptide epitope of the polypeptide of claim 3.
10. The polypeptide of claim 3 wherein the polypeptide has biological activity.
11. An isolated nucleic acid encoding a species homologue of the polypeptide of claim 3.
12. The isolated nucleic acid molecule of claim 1, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the 5' end or the 3' end.
13. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.
14. A recombinant host cell comprising the isolated nucleic acid molecule of claim 1.

15. A method of making the recombinant host cell of claim 14 comprising transforming a host cell with the nucleic acid molecule of claim 1.
16. The recombinant host cell of claim 14 comprising vector sequences.
17. The isolated polypeptide of claim 3, wherein the isolated polypeptide
5 comprises sequential amino acid deletions from either the C-terminus or the N-terminus.
18. An isolated antibody that binds specifically to the isolated polypeptide of claim 3.
19. The isolated antibody of claim 18 wherein the antibody is a monoclonal antibody.
- 10 20. The isolated antibody of claim 18 wherein the antibody is a polyclonal antibody.
21. A recombinant host cell that expresses the isolated polypeptide of claim 3.
22. An isolated polypeptide produced by the steps of:
(a) culturing the recombinant host cell of claim 14 under conditions such
15 that said polypeptide is expressed; and
(b) isolating the polypeptide.
23. A method for preventing, treating, modulating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 3 or the polynucleotide of claim 1.
- 20 24. The method of claim 23 wherein the medical condition is inflammatory bowel disease.
25. A method for preventing, treating, modulating, or ameliorating a medical condition comprising administering to a mammalian subject a therapeutically effective amount of the antibody of claim 18.
- 25 26. The method of claim 25 wherein the medical condition is inflammatory bowel disease.
27. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
(a) determining the presence or absence of a mutation in the polynucleotide of
30 claim 1; and
(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.

28. The method of claim 27 wherein the pathological condition is inflammatory bowel disease.

29. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising detecting an alteration in expression of a polypeptide of claim 3, wherein the presence of an alteration in expression of the polypeptide is indicative of the pathological condition or susceptibility to the pathological condition.

30. The method of claim 29 wherein the alteration in expression is an increase in the amount of expression or a decrease in the amount of expression.

31. The method of claim 29 wherein the pathological condition is inflammatory bowel disease.

32. The method of claim 31 wherein the method further comprises the steps of: obtaining a first biological sample from a patient suspected of having inflammatory bowel disease and obtaining a second sample from a suitable comparable control source;

(a) determining the amount of at least one polypeptide encoded by a polynucleotide of claim 1 in the first and second sample; and
(b) comparing the amount of the polypeptide in the first and second samples; wherein a patient is diagnosed as having inflammatory bowel disease if the amount of the polypeptide in the first sample is greater than or less than the amount of the polypeptide in the second sample.

33. The use of the polynucleotide of claim 1 or polypeptide of claim 3 for the manufacture of a medicament for the treatment of inflammatory bowel disease.

34. The use of the antibody of claim 18 for the manufacture of a medicament for the treatment of inflammatory bowel disease.

35. A method for identifying a binding partner to the polypeptide of claim 3 comprising:

(a) contacting the polypeptide of claim 3 with a binding partner; and
(b) determining whether the binding partner effects an activity of the polypeptide.

36. The gene corresponding to the cDNA sequence of the isolated nucleic acid of claim 1.

37. A method of identifying an activity of an expressed polypeptide in a biological assay, wherein the method comprises:

- (a) expressing the polypeptide of claim 3 in a cell;
- (b) isolating the expressed polypeptide;
- (c) testing the expressed polypeptide for an activity in a biological assay; and
- (d) identifying the activity of the expressed polypeptide based on the test results.

5 38. A substantially pure isolated DNA molecule suitable for use as a probe for genes regulated in inflammatory bowel disease, chosen from the group consisting of the DNA molecules identified in Table 1, having a 5' partial nucleotide sequence and length as described by their digital address, and having a characteristic regulation pattern in inflammatory bowel disease.

10 39. A kit for detecting the presence of the polypeptide of the claim 3 in a mammalian tissue sample comprising a first antibody which immunoreacts with a mammalian protein encoded by a gene corresponding to the polynucleotide of claim 1 or with a polypeptide encoded by the polynucleotide of claim 3 in an amount sufficient for at least one assay and suitable packaging material.

15 40. A kit of claim 39 further comprising a second antibody that binds to the first antibody.

 41. The kit of claim 40 wherein the second antibody is labeled.

 42. The kit of claim 41 wherein the label comprises enzymes, radioisotopes, fluorescent compounds, colloidal metals, chemiluminescent compounds, phosphorescent
20 compounds, or bioluminescent compounds.

 43. A kit for detecting the presence of a genes encoding an protein comprising a polynucleotide of claim 1, or fragment thereof having at least 10 contiguous bases, in an amount sufficient for at least one assay, and suitable packaging material.

 44. A method for detecting the presence of a nucleic acid encoding a protein in a
25 mammalian tissue sample, comprising the steps of:

- (a) hybridizing a polynucleotide of claim 1 or fragment thereof having at least 10 contiguous bases, with the nucleic acid of the sample; and
- (b) detecting the presence of the hybridization product.

30

ABSTRACT OF THE DISCLOSURE

The invention is directed to purified and isolated novel polypeptides, the nucleic acids encoding such polypeptides, processes for production of recombinant forms of such polypeptides, antibodies generated against these polypeptides, fragmented peptides derived
5 from these polypeptides, and the uses of the above.